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Evolution of the hypoxia-sensitive cells involved in amniote respiratory reflexes

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Abstract The evolutionary origins of the hypoxia-sensitive cells that trigger amniote respiratory reflexes – carotid body glomus cells, and ‘pulmonary neuroendocrine cells’ (PNECs) – are obscure. Homology has been proposed between glomus cells, which are neural crest-derived, and the hypoxia-sensitive ‘neuroepithelial cells’ (NECs) of fish gills, whose embryonic origin is unknown. NECs have also been likened to PNECs, which differentiate in situ within lung airway epithelia. Using genetic lineage-tracing and neural crest-deficient mutants in zebrafish, and physical fate-mapping in frog and lamprey, we find that NECs are not neural crest-derived, but endoderm-derived, like PNECs, whose endodermal origin we confirm. We discover neural crest-derived catecholaminergic cells associated with zebrafish pharyngeal arch blood vessels, and propose a new model for amniote hypoxia-sensitive cell evolution: endoderm-derived NECs were retained as PNECs, while the carotid body evolved via the aggregation of neural crest-derived catecholaminergic (chromaffin) cells already associated with blood vessels in anamniote pharyngeal arches.

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eLife digest The carotid bodies are small glands found in either side of our neck, near the carotid artery. When the level of oxygen in our blood drops, specialized cells in the carotid bodies signal to the brain to increase our heart rate and make us breathe more rapidly and deeply. As a result, more oxygen is delivered to our cells.

Fish have similar oxygen-sensitive cells in their gills, known as neuroepithelial cells, that detect changes in the oxygen levels in the surrounding water and their blood. It has been suggested that after our vertebrate (back-boned animal) ancestors moved onto land, the neuroepithelial cells in their gills eventually evolved to form the carotid bodies. Knowing whether this is true would allow researchers to better understand how our ancestors were able to adapt to an obligate air-breathing lifestyle on land.

If the carotid body did evolve from ancestral neuroepithelial cells, we would expect that they would both develop from the same kind of cells in the embryo. Carotid body cells develop from a group of cells called neural crest cells, which give rise to many tissues, including nerve cells. Hockman et al. have now investigated whether neuroepithelial cells also develop from neural crest cells.

Hockman et al. labelled the neural crest cells in the embryos of zebrafish, frogs and lampreys using techniques such as injecting the cells with fluorescent dye or genetically modifying the cells to make fluorescent proteins. Unexpectedly, the neuroepithelial cells that developed in the gills of these embryos did not contain these fluorescent labels, meaning that they did not develop from the neural crest cells. The patterns of gene activity found in the developing neuroepithelial cells were also different from those in the carotid body. Further investigation revealed that neuroepithelial cells develop from the lining of the mouth and gills and may be related to a similar population of oxygen-sensitive cells found in the lungs.

Overall, it appears that the carotid body did not evolve from ancestral neuroepithelial cells. However, Hockman et al. did find some cells near blood vessels in the gills of zebrafish that had developed from neural crest cells. Equivalent cells in our ancestors could therefore be the cells that evolved into carotid bodies. A first test of this theory will be to determine whether or not these cells are oxygen-sensitive.

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Introduction

During hypoxia in vertebrates, respiratory reflexes such as hyperventilation are triggered by neurotransmitter release from hypoxia-sensitive serotonergic cells associated with pharyngeal arch arteries, as well as in the lungs and/or gills (reviewed by [López-Barneo et al., 2016](#); [Cutz et al., 2013](#); [Jonz et al., 2016](#)). In amniotes, these are the ‘glomus cells’ of the carotid body, located at the bifurcation of the common carotid artery (reviewed by [Nurse, 2014](#); [López-Barneo et al., 2016](#)), and the ‘pulmonary neuroendocrine cells’ (PNECs) of lung airway epithelia, found either as solitary, flask-shaped cells, or collected into ‘neuroepithelial bodies’ (NEBs), preferentially located at airway branch points (reviewed by [Cutz et al., 2013](#)). Glomus cells respond to hypoxia in arterial blood by releasing stored neurotransmitters including acetylcholine, ATP, the catecholamine dopamine, and serotonin (the latter two most likely acting as autocrine/paracrine neuromodulators) (reviewed by [Nurse and Piskuric, 2013](#); [Nurse, 2014](#)). These excite afferent terminals of the carotid sinus nerve (a branch of the glossopharyngeal nerve, arising from neurons in the petrosal ganglion) in mammals (reviewed by [Nurse and Piskuric, 2013](#); [Nurse, 2014](#)), and of the vagal nerve (arising from neurons in the nodose ganglion) in birds ([Kameda, 2002](#)). The afferent nerves relay signals to the nucleus of the solitary tract within the hindbrain, to elicit respiratory reflex responses such as hyperventilation (reviewed by [Teppema and Dahan, 2010](#)). PNECs respond to hypoxia by releasing stored serotonin and various neuropeptides onto vagal afferents, and are thought to act as hypoxia-sensitive airway sensors (reviewed by [Cutz et al., 2013](#); also see [Branchfield et al., 2016](#)). PNECs also provide an important stem-cell niche for regenerating the airway epithelium after injury ([Reynolds et al., 2000](#); [Guha et al., 2012](#); [Song et al., 2012](#)) and were recently shown to be the predominant cells of origin for small cell lung cancer ([Park et al., 2011](#); [Sutherland et al., 2011](#); [Song et al., 2012](#)).

The evolution of glomus cells was critical for the transition from aquatic life, where externally facing hypoxia-sensors are essential for monitoring the variable oxygen levels in water, to fully terrestrial life, where reflex responses to variations in internal oxygen levels are more important, given the stability of oxygen levels in air (Burleson and Milsom, 2003; Milsom and Burleson, 2007). However, the evolutionary history of glomus cells - and, indeed, PNECs - is uncertain. One commonly suggested hypothesis (e.g., Milsom and Burleson, 2007; Hempleman and Warburton, 2013; Jonz et al., 2016) is that glomus cells are homologous to the chemosensory 'neuroepithelial cells' (NECs) of fish gills. These were originally identified within the primary epithelium of the gills in various teleosts and a shark, as innervated cells (isolated or clustered) containing dense-cored vesicles; formaldehyde-induced fluorescence revealed the presence of biogenic amines, identified as serotonin, while electron microscopy following exposure of trout to acute hypoxia revealed fewer vesicles, which appeared degranulated (Dunel-Erb et al., 1982). In vitro patch-clamp studies on NECs isolated from zebrafish and catfish gills (identified by neutral red, a vital dye that stains monoamine-containing cells including serotonergic PNECs; Youngson et al., 1993), confirmed that teleost gill NECs are hypoxia-sensitive (Jonz et al., 2004; Burleson et al., 2006). The conventional marker for teleost gill NECs is serotonin: although non-serotonergic gill NECs were identified in adult zebrafish by immunoreactivity for synaptic vesicle glycoprotein 2, these may represent immature NECs (Jonz and Nurse, 2003; Jonz et al., 2004). NECs are found near efferent gill arteries and on the basal lamina of the gill epithelia, facing the flow of water, hence can detect hypoxia and other stimuli in either blood or external water (reviewed by Jonz et al., 2016). Like glomus cells (reviewed by Nurse, 2014; López-Barneo et al., 2016), zebrafish gill NECs also respond to acid hypercapnia (increased CO_2/H^+) (López-Barneo et al., 1988; Buckler, 1997; Jonz et al., 2004; Qin et al., 2010). The putative shared evolutionary ancestry of glomus cells and gill NECs (e.g. Milsom and Burleson, 2007; Hempleman and Warburton, 2013; Jonz et al., 2016) is supported by many similarities: both are associated with pharyngeal arch arteries (the carotid body develops in association with the third pharyngeal arch artery, which will form the carotid artery), provided with afferent innervation by glossopharyngeal and/or vagal nerves, and have background ('leak') K^+ currents that are inhibited by hypoxia, resulting in membrane depolarization, activation of voltage-gated Ca^{2+} channels, and neurotransmitter release (López-Barneo et al., 1988; Buckler, 1997; Jonz et al., 2004; Qin et al., 2010).

If glomus cells and gill NECs evolved from the same ancestral cell population, they should share a common embryonic origin. Glomus cells are neural crest-derived, as demonstrated in birds by quail-chick neural fold grafts (Le Douarin et al., 1972; Pearse et al., 1973), and in mouse by *Wnt1-Cre* genetic lineage-tracing (Pardal et al., 2007), but the embryonic origin of NECs is unknown. Lack of immunoreactivity for the HNK1 antibody, which labels migrating neural crest cells in many but not all vertebrates, has been reported for NECs (Porteus et al., 2013, 2014). However, the carbohydrate epitope recognized by the HNK1 antibody (Voshol et al., 1996) is borne by multiple glycoproteins and glycolipids, and gene or antigen expression in itself cannot indicate lineage. An alternative to the hypothesis that gill NECs and glomus cells evolved from a common ancestral cell population is that NECs share ancestry with PNECs, to which they were originally likened (Dunel-Erb et al., 1982). Hypoxia-sensing by PNECs, as in NECs and glomus cells, involves inhibition of a K^+ current by hypoxia (reviewed by Cutz et al., 2013; Nurse, 2014; López-Barneo et al., 2016; Jonz et al., 2016). In contrast to the neural crest origin of glomus cells (Le Douarin et al., 1972; Pearse et al., 1973; Pardal et al., 2007), PNECs have an intrinsic pulmonary epithelial origin: the first experimental support for this was provided by a tritiated thymidine labeling study of hamster lung development (Hoyt et al., 1990), later confirmed by mouse genetic lineage-tracing studies using *Id2-CreER^{T2}*, *Shh-Cre*, *Nkx2.1-Cre*, and *Sox9-Cre* driver lines that showed a common origin for all lung airway epithelial cell types, including PNECs (Rawlins et al., 2009; Song et al., 2012; Kuo and Krasnow, 2015).

Here, we demonstrate that neural crest cells do not contribute to gill NECs: instead, these are endoderm-derived. This refutes the hypothesis that glomus cells and gill NECs evolved from a common ancestral cell population, and instead supports an evolutionary relationship between NECs and PNECs, whose endodermal origin we confirm in mouse. We also show that the transcription factor *Phox2b*, which is required for glomus cell development (Dauger et al., 2003), is not expressed by gill NECs (or by PNECs), arguing against the possibility of cell-type homology between glomus cells and gill NECs via activation of the same genetic network. Finally, we report the discovery of neural

crest-derived chromaffin (catecholaminergic) cells associated with blood vessels in the pharyngeal arches of juvenile zebrafish, which we speculate could share an evolutionary ancestry with glomus cells. Given these results, we propose a new model for the evolution of hypoxia-sensitive cells during the transition to terrestrial life.

Results

Carotid body glomus cells develop from the neural crest (Le Douarin et al., 1972; Pearse et al., 1973; Pardal et al., 2007), while PNECs differentiate in situ within pulmonary airway epithelia (Hoyt et al., 1990; Rawlins et al., 2009; Song et al., 2012; Kuo and Krasnow, 2015). We aimed to shed light on the evolutionary origins of these amniote hypoxia-sensitive cell types by determining the embryonic origin of NECs, the hypoxia-sensitive cells of anamniote gills.

Zebrafish NECs are not neural crest-derived

In developing zebrafish, gill NECs were previously identified as serotonin (5-HT)-immunoreactive cells in gill filaments from 5-dpf, which are innervated by 7-dpf (Jonz and Nurse, 2005). We investigated any neural crest contribution to gill NECs via genetic lineage-tracing, using a collection of transgenic zebrafish lines with different *cis*-regulatory sequences driving Cre and subsequent lineage reporter expression in neural crest-derived cells, as well as via neural crest-deficient zebrafish embryos. Lineage-tracing using different Cre driver and reporter lines enabled us to control for false negatives potentially arising from variable promoter activity in, or incomplete labeling of, some neural crest cells in either neural crest driver or reporter lines.

In larval and juvenile zebrafish, we identified NECs in the gill filaments by serotonin immunoreactivity, as previously reported (Jonz and Nurse, 2005); we also found similar innervated serotonergic cells scattered in the orobranchial epithelium (putative NECs). NECs in the gill filaments, and putative NECs in the orobranchial epithelium, were unlabeled in larvae/metamorphic juveniles with *sox10* *cis*-regulatory sequences driving Cre and resulting lineage reporter expression [*Tg(-28.5sox10:cre); Tg(ef1a:loxP-DsRed-loxP-EGFP)* (Kague et al., 2012); *Tg(-4.9sox10:creER^{T2}); Tg(βactin:loxP-Super-Stop-loxP-DsRed)* (Mongera et al., 2013)], even when nearby neural crest derivatives such as branchial arch cartilages/mesenchyme and/or gill pillar cells (Mongera et al., 2013) were reporter-positive (Figure 1a–f'; 183 serotonergic cells located near such reporter-positive cells [≥ 6 per fish] were counted across 11 larvae/metamorphic juveniles: $n = 8$ for *-28.5sox10*; $n = 3$ for *-4.9sox10*). Gill NECs, and putative NECs in the orobranchial epithelium, were also unlabeled in *Tg(crestin:creER^{T2}); Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* (Mosimann et al., 2011; Kaufman et al., 2016) larvae/metamorphic juveniles, even when nearby neural crest-derived cells in branchial arch cartilages and/or gill filament mesenchyme were mCherry-positive (Figure 1g–h'; 166 serotonergic cells located near such mCherry-positive cells [≥ 6 per fish] were counted across 10 larvae/metamorphic juveniles). We ruled out the possibility that lack of lineage reporter expression in NECs was a false-negative result arising from inactivity in NECs of the promoters driving the Cre-switchable reporter cassettes, by confirming that NECs expressed the native unrecombined reporter in both transgenic lines [*Tg(-28.5sox10:cre); Tg(ef1a:loxP-DsRed-loxP-EGFP)* and *Tg(crestin:creER^{T2}); Tg(-3.5ubi:loxP-GFP-loxP-mCherry)*] (Figure 1—figure supplement 1).

Finally, we analyzed *tfap2a^{mob}; foxd3^{mos}* zebrafish, which lack neural crest derivatives (Barralho-Gimeno et al., 2004; Montero-Balaguer et al., 2006; Wang et al., 2011). At 7-dpf, *tfap2a^{mob}; foxd3^{mos}* mutants lacked pigment cells and lower jaw structures ($n = 8$; Figure 2a). In the absence of the neural crest-derived pharyngeal endoskeleton, pharyngeal arches and gills are hard to recognize, but putative NECs, visualized here as serotonergic cells associated with HNK1 epitope-immunoreactive neurites (Metcalf et al., 1990), were still present in the orobranchial epithelium (Figure 2b,c) (and occasionally could also be identified ventral to the orobranchial cavity, where the pharyngeal arches would be located; Figure 2d,e). We counted all putative NECs in the orobranchial region of three *tfap2a^{mob}; foxd3^{mos}* larvae (294 putative NECs counted in total) and three wild-type siblings (244 putative NECs counted in total): there was no change in mean number (mean/embryo \pm s.d.: 81.3 ± 24.0 for wild-type larvae, $n = 3$; 98.0 ± 49.7 for *tfap2a^{mob}; foxd3^{mos}* larvae, $n = 3$; $p = 0.63$, unpaired two-tailed Student's t-test) (Figure 2f–i).

Putative NECs have also been identified as serotonergic cells in the skin of embryonic zebrafish (Jonz and Nurse, 2006; Coccimiglio and Jonz, 2012) and of adult mangrove killifish, which respire

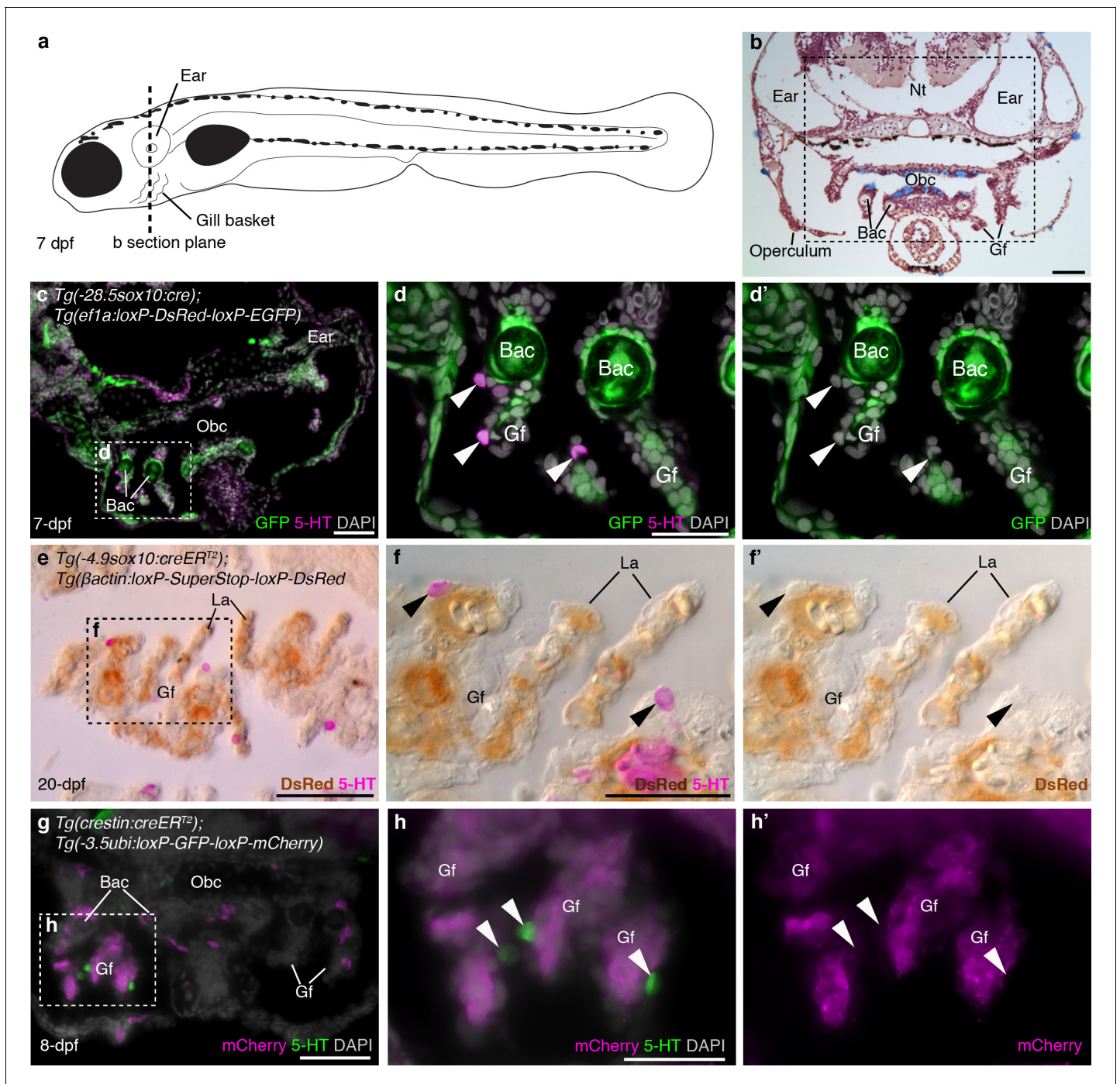


Figure 1. Zebrafish NECs are not neural crest-derived: genetic lineage-tracing data. (a) Schematic of a 7–8 dpf zebrafish; dotted line indicates section plane in b–f'. (b) Hematoxylin and eosin staining at 7-dpf reveals gill filaments branching from branchial arch cartilages, and the orobranchial cavity. Dashed box indicates approximate region in c,g. (c–d') In 7-dpf *Tg(-28.5sox10:cre);Tg(ef1a:loxP-DsRed-loxP-EGFP)* zebrafish, GFP labels neural crest-derived branchial arch cartilage and mesenchyme, but not NECs in the gill filaments (identified by immunoreactivity for serotonin, 5-HT; arrowheads). (e–f') Horizontal section through the gills of a 20-dpf *Tg(-4.9sox10:creER^{T2});Tg(beta1a:loxP-SuperStop-loxP-DsRed)* zebrafish. DsRed (brown precipitate) labels neural crest-derived gill pillar cells, but not NECs (arrowheads; inverted fluorescent image overlaid on bright-field image). (g–h') In 8-dpf *Tg(crestin:creER^{T2});Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* zebrafish, mCherry labels gill pillar cells but not NECs (arrowheads). 5-HT, serotonin; Bac, branchial arch cartilage; Gf, gill filament; La, lamellae; Nt, neural tube; Obc, orobranchial cavity. Scale-bars: 50 μ m in b,c,e,g; 25 μ m in d,f,h.

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The following figure supplement is available for figure 1:

Figure supplement 1. The promoters driving the Cre-switchable reporter cassettes are active in NECs.

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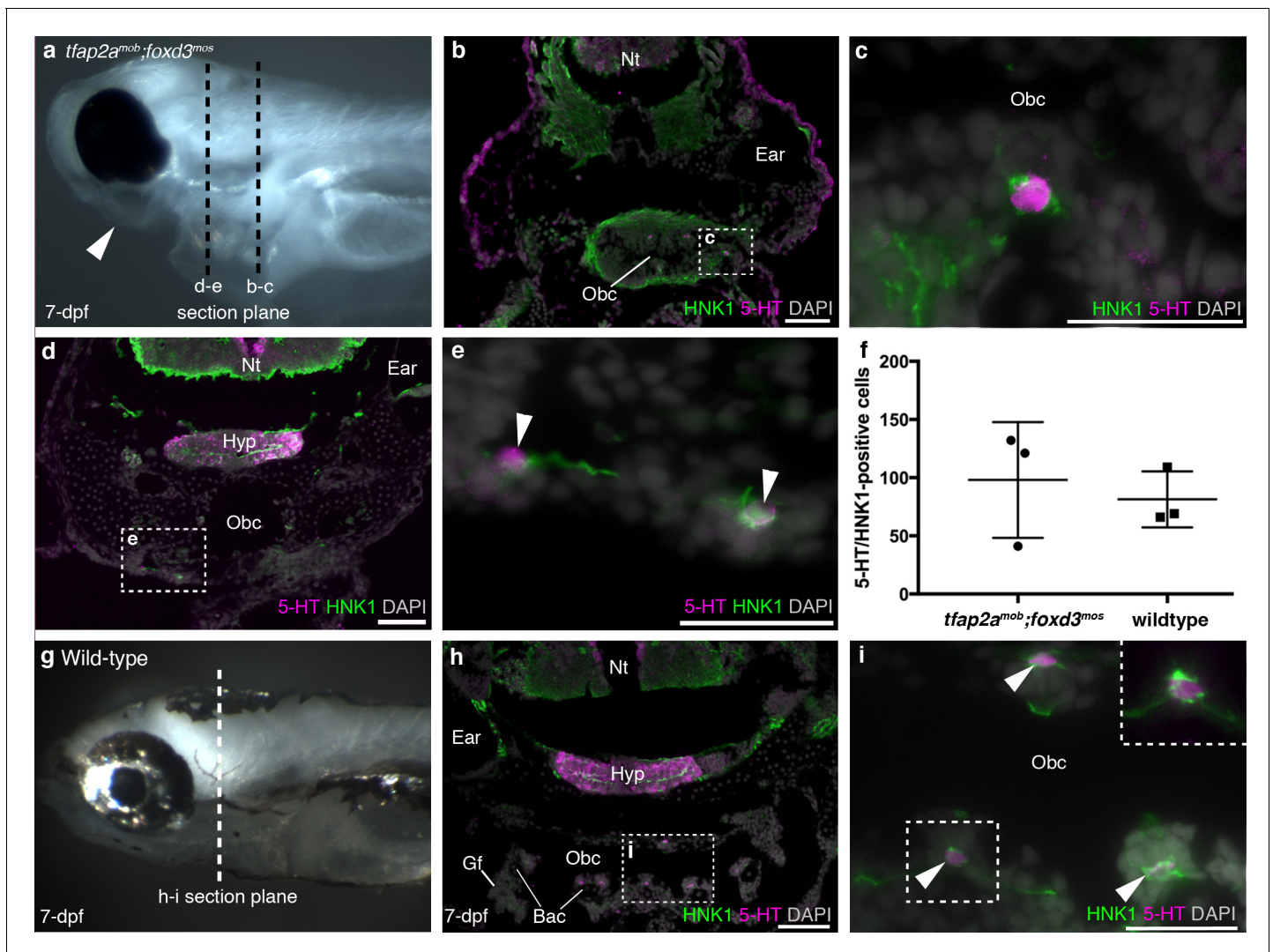


Figure 2. Zebrafish NECs are not neural crest-derived: analysis of neural crest-deficient zebrafish mutants. (a) 7-dpf *tfap2a^{mob};foxd3^{mos}* zebrafish lack all neural crest derivatives, including melanophores and jaw skeleton (arrowhead). Dotted lines: section planes in b-e. (b,c) At 7-dpf, *tfap2a^{mob};foxd3^{mos}* orobranchial epithelium retains innervated serotonergic (5-HT⁺) cells (putative NECs), identified by cytoplasmic serotonin surrounded by a ring of HNK1 epitope-immunoreactive neurites. (d,e) At 7-dpf, putative NECs (arrowheads) persist in *tfap2a^{mob};foxd3^{mos}* zebrafish in the region ventral to the orobranchial cavity where the pharyngeal arches would be located in wild-type fish. (NB The hypothalamus [Hyp] extends caudally beneath the midbrain and rostral hindbrain, and often separates from the overlying brain on sections, as here.) (f) The mean number per 7-dpf larva of putative NECs in the orobranchial epithelium does not differ between *tfap2a^{mob};foxd3^{mos}* (98.0 ± 49.7 s.d.; $n = 3$) and wild-type zebrafish (81.3 ± 24.0 s.d.; $n = 3$) ($p=0.63$, unpaired two-tailed Student's t-test). All such cells in the orobranchial epithelium were counted for each embryo. Error bars indicate s.d. (g-i) Wild-type sibling at 7-dpf. Dotted line: section plane in h-i. Putative NECs are present in the orobranchial epithelium (arrowheads; dashed box in i, magnified without DAPI in top right corner). 5-HT, serotonin; Gf, gill filament; Hyp, hypothalamus; Nt, neural tube; Obc, orobranchial cavity. Scale-bars: 50 μ m in b,d,h; 25 μ m in c,e,i.

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The following figure supplement is available for figure 2:

Figure supplement 1. Putative NECs in the skin of embryonic zebrafish are not neural crest-derived.

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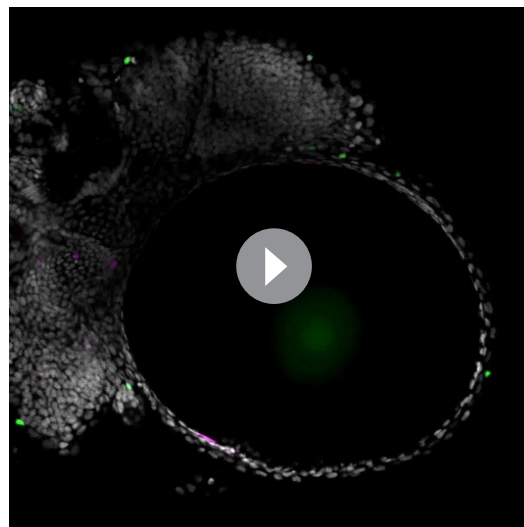
through the skin as well as the gills (Regan et al., 2011). Although they have not been shown directly (e.g., by patch-clamp experiments) to be hypoxia-sensitive, the putative NECs in killifish increase in area in response to hypoxia (Regan et al., 2011), while in zebrafish, hypoxia decreased or delayed, and hyperoxia accelerated, the normal decline in number of these cells seen with increasing age (Coccimiglio and Jonz, 2012). In zebrafish, these serotonergic cells are most

abundant at 3-dpf, and most evident scattered in the skin over the eyes, yolk-sac and tail (**Coccimiglio and Jonz, 2012**). Whole-mount immunostaining of *Tg(crestin:creER^{T2});Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* embryos at 3-dpf for serotonin and mCherry revealed the expected pattern of scattered serotonergic cells in the epidermis, but none was mCherry-positive (i.e., neural crest-derived) (**Figure 2—figure supplement 1** shows a sample three-dimensional rendering for the eye; for the associated z-stack movies, see **Videos 1** and **2**). We quantified this in the eye, where the corneal endothelium is neural crest-derived, as previously reported for chicken (**Noden, 1978; Johnston et al., 1979**), mouse (with a minor mesodermal contribution also noted; **Gage et al., 2005**) and *Xenopus* (**Hu et al., 2013**). At 3-dpf, the zebrafish cornea comprises an outer corneal epithelium, a thin acellular collagenous stroma, and a monolayer of flattened corneal endothelial cells (**Soules and Link, 2005; Zhao et al., 2006; Akhtar et al., 2008**). Across 13 embryos, we counted 328 serotonergic cells in the epidermis over the eye, all of which were mCherry-negative, and 219 nearby mCherry-positive (i.e., neural crest-derived) corneal endothelial cells. Thus, the putative NECs in the skin of embryonic zebrafish are not neural crest-derived.

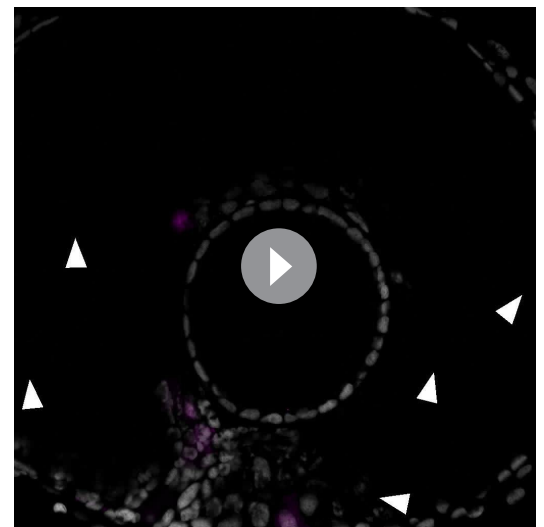
Taken together, these results show that in zebrafish (a ray-finned fish), the neural crest does not contribute to gill NECs (the previously proposed homologs of glomus cells), or to putative NECs in the orobranchial epithelium and embryonic epidermis.

Putative NECs in *Xenopus* and lamprey are not neural crest-derived

Innervated serotonergic cells in the internal gills of *Xenopus* tadpoles are the proposed homologs of teleost gill NECs (**Saltys et al., 2006**). We detected scattered serotonergic cells in gill filaments from stage 43 (**Figure 3a,b**), and in the orobranchial epithelium from stage 41 (**Figure 3a,c**). To determine if these putative NECs are neural crest-derived, we unilaterally grafted neural folds from GFP-labeled donors to unlabeled hosts (**Figure 3d**). GFP-labeled neural crest cells migrated away



Video 1. Putative NECs in the skin of embryonic zebrafish are not neural crest-derived. Z-stack movie showing a whole-mount view of the eye region of a 3-dpf *Tg(crestin:creER^{T2});Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* embryo, immunostained for serotonin and mCherry. Serotonergic cells are scattered in the epidermis over the eye, as reported (**Coccimiglio and Jonz, 2012**), but none is mCherry-positive, i.e., neural crest-derived. **Video 2** shows a higher-power movie. A three-dimensional rendering for the eye is shown in **Figure 2—figure supplement 1**. DOI: 10.7554/eLife.21231.007



Video 2. Putative NECs in the skin of embryonic zebrafish are not neural crest-derived. Z-stack movie at higher power than **Video 1**, showing a whole-mount view of the eye region of a 3-dpf *Tg(crestin:creER^{T2});Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* embryo, immunostained for serotonin and mCherry. Serotonergic cells (arrowheads) are scattered in the epidermis over the eye, as reported (**Coccimiglio and Jonz, 2012**), but none is mCherry-positive, i.e., neural crest-derived. A 3-dimensional rendering for the eye is shown in **Figure 2—figure supplement 1**. DOI: 10.7554/eLife.21231.008

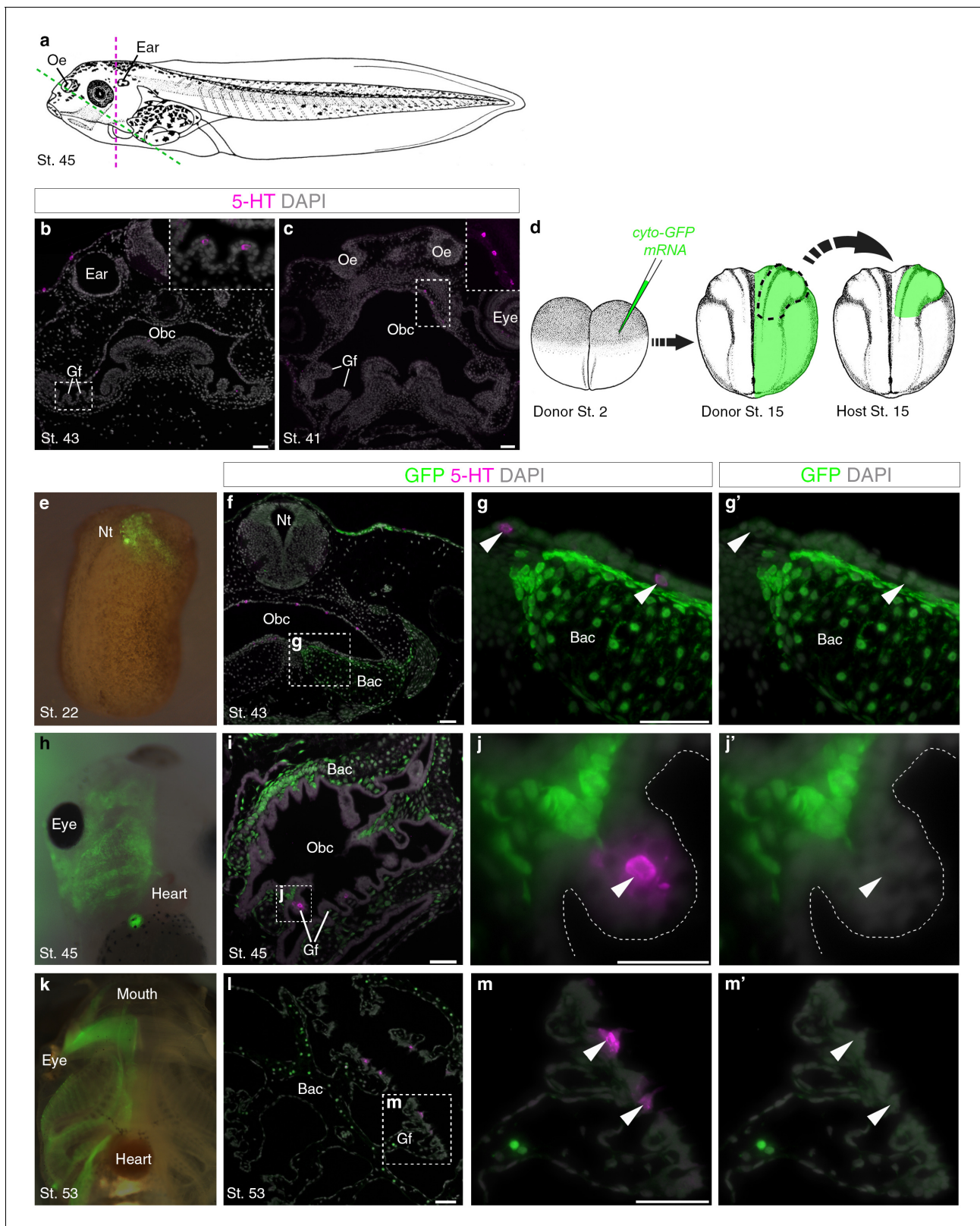


Figure 3. Putative *Xenopus* NECs are not neural crest-derived. (a) Schematic of a stage 45 tadpole (*Nieuwkoop and Faber, 1967*). Dotted lines: section planes in **b,f–g'** (magenta, transverse) and in **c,i–j',l–m'** (green, oblique). (b) Serotonergic cells in gill filament epithelium (putative NECs) are first detected at stage 43 (inset shows higher-power view). (c) Similar serotonergic cells are scattered in the orobranchial epithelium from stage 41 (inset shows higher-power view). (d) Schematic (modified from *Nieuwkoop and Faber, 1967*) showing neural crest labelling: GFP-donors, created by Figure 3 continued on next page

Figure 3 continued

injecting *cyto-GFP* mRNA into one cell at the two-cell stage, were grown to stages 13–17, and neural folds grafted unilaterally to wild-type hosts. For grafted embryos grown to stage 53, donors were transgenic *CMV-GFP* embryos. (e) At stage 22, GFP labels neural crest cells migrating towards the branchial arches. (f–g) At stage 43, GFP labels branchial arch cartilage and surrounding mesenchyme, but not putative NECs (arrowheads) in the orobranchial epithelium. (h–j) At stage 45, GFP-positive neural crest cells are visible in the branchial arches (whole-mount and section from different embryos). GFP labels branchial arch cartilage and mesenchyme, but not putative NECs (arrowheads) in the gill filaments. (k–m) At stage 53 (transgenic *CMV-GFP* donors; whole-mount and section from different embryos), GFP labels branchial arch cartilage and mesenchyme, but not putative NECs (arrowheads) in the gill filaments. 5-HT, serotonin; Bac, branchial arch cartilage; Gf, gill filament; Nt, neural tube; Obc, orobranchial cavity; Oe, olfactory epithelium. Scale-bar: 50 μ m.

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from the neural tube and contributed to branchial arch cartilages and mesenchyme, but putative NECs in the gill filaments and orobranchial epithelium were GFP-negative ($n = 14$; **Figure 3e–m**).

In the sea lamprey, the gills are arranged in pairs within the orobranchial cavity, supported by an interbranchial septum (**Figure 4a–c**). Serotonergic cells in lamprey gills are proposed to correspond to the gill NECs of jawed fishes (**Barreiro-Iglesias et al., 2009**). We detected putative NECs from embryonic day (E)18.5, in clusters on the medial edges of the gills, intimately associated with HNK1 epitope-immunoreactive neurites (**Figure 4d**). Scattered serotonergic cells were also found in the epithelium lining the roof and floor of the orobranchial cavity. To determine any neural crest cell contribution, the vital lipophilic dye Dil was injected into E5 vagal neural folds and the embryos followed to E19.0 (**Figure 4e–n**). Dil-labeled neural crest cells migrated into the branchial arches and contributed to the branchial arch basket, as expected (**McCauley and Bronner-Fraser, 2003**), but putative NECs in the gills and orobranchial epithelia, despite being near Dil-labeled cells, were unlabeled ($n = 19$; **Figure 4i,j,n**; **Figure 4—figure supplement 1**).

These results show that putative NECs in the internal gills and orobranchial epithelium of a frog (i.e., a lobe-finned tetrapod) and the sea lamprey (a jawless fish) are not neural crest-derived.

Gill NECs are endoderm-derived, like PNECs

Overall, our data show that the neural crest does not contribute to gill NECs in zebrafish, or to their presumed homologues in *Xenopus* and lamprey gills (or to the putative NECs identified in the orobranchial epithelium of all three species). Hence, glomus cells and gill NECs cannot have evolved from the same ancestral cell population. In zebrafish, *Xenopus* and the little skate (a cartilaginous fish), vital dye fate-mapping experiments have shown that the gills and orobranchial cavity are lined with an epithelium derived mostly from endoderm (**Warga and Nüsslein-Volhard, 1999; Chalmers and Slack, 2000; Gillis and Tidswell, 2017**), suggesting endoderm as an alternative origin for NECs. Indeed, in one of the first descriptions of fish gill NECs (**Dunel-Erb et al., 1982**), they were likened to the PNECs of amniotes, which share a common embryonic origin with other airway epithelial cells in rodents (**Hoyt et al., 1990; Rawlins et al., 2009; Song et al., 2012; Kuo and Krasnow, 2015**). We demonstrated the endodermal origin of PNECs in the mouse by lineage-tracing using the *Sox17*^{2A-iCre} driver line (in which all endoderm-derived lineages, as well as vascular endothelial cells and the hematopoietic system, express Cre; **Engert et al., 2009**) crossed to the *R26R*^{lacZ} or *R26R*^{tdTomato} reporter lines (**Soriano, 1999; Madisen et al., 2010**) (**Figure 5a–e''**). We also confirmed the recent exclusion by *Wnt1*-Cre lineage-tracing (**Danielian et al., 1998**) of a neural crest contribution to mouse PNECs (**Kuo and Krasnow, 2015**) (**Figure 5—figure supplement 1a–b'**). Similarly, we saw no neural crest contribution to PNECs in the chicken lung after labeling the premigratory neural crest by neural fold grafting from GFP-transgenic donor embryos (**McGrew et al., 2008**) (**Figure 5—figure supplement 1c–e'**).

To test the hypothesis that NECs, like PNECs, are endoderm-derived, we first attempted to label the pharyngeal endoderm of *Xenopus* embryos at stage 14 via focal Dil injections into anterior endoderm (**Chalmers and Slack, 2000**) (**Figure 5f–i**). Only three embryos with endoderm-specific Dil labeling survived to stage 45 for analysis. The Dil labeling was very sparse, but in one embryo, 15 serotonergic cells in the orobranchial epithelium (putative NECs) were Dil-labeled, supporting an endodermal origin (**Figure 5g–i''**).

Since the direct labeling approach in *Xenopus* proved to be technically challenging, we used genetic lineage-tracing of endodermal *sox17* expression in zebrafish, inducing gastrulation-stage

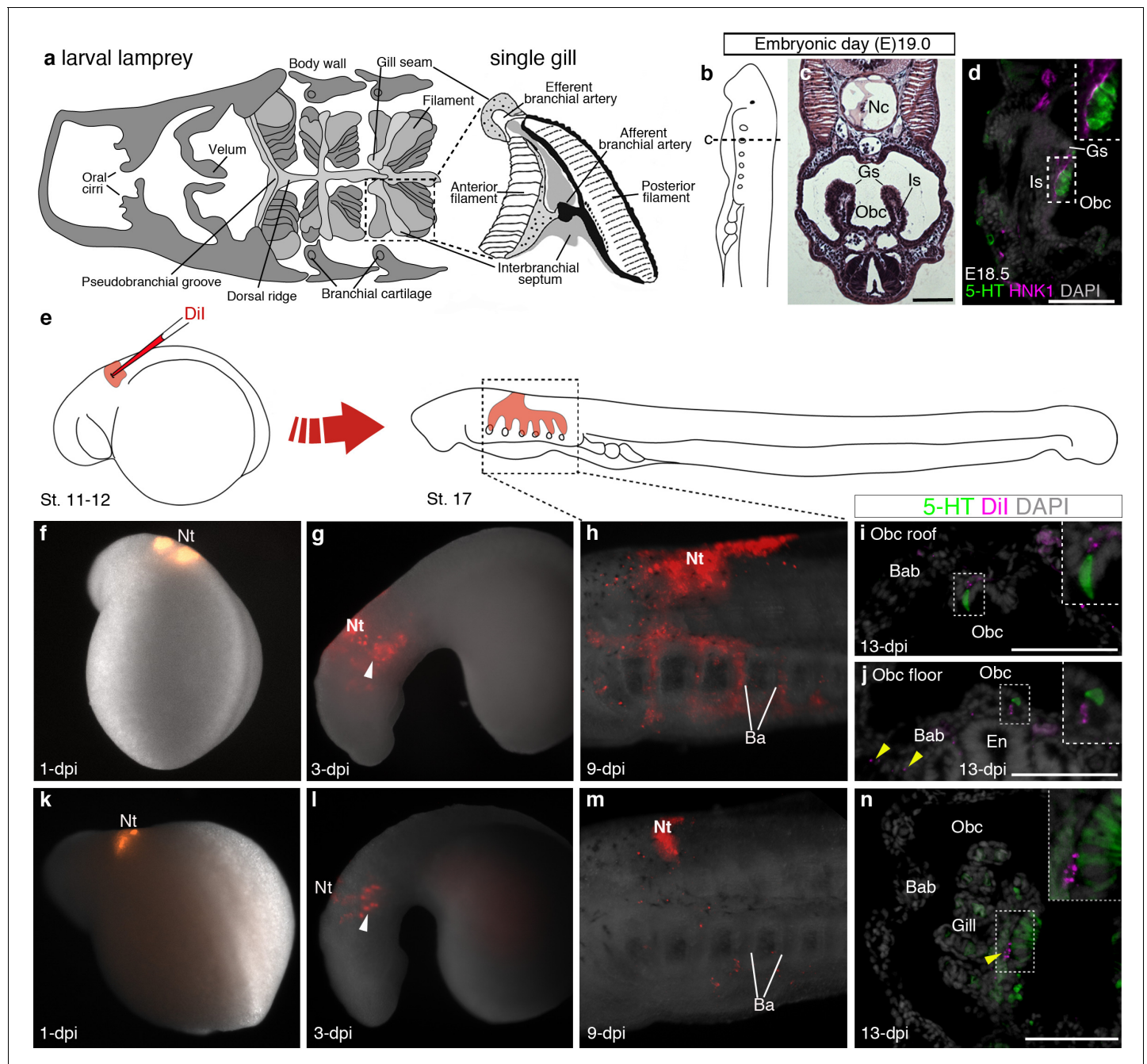


Figure 4. Putative lamprey NECs are not neural crest-derived. (a) Schematic larval lamprey section (modified from [Barreiro-Iglesias et al., 2009](#)) showing gill pairs in the orobranchial cavity, supported by an interbranchial septum, and a single gill at higher power. (b) Schematic Piavis-stage 17 (E19) lamprey (modified from [Tahara, 1988](#)). Dotted line shows section plane in **c,d,i,j,n**. (c) Hematoxylin and eosin staining at E19 shows internal gills as 'stalks' within the orobranchial cavity. (d) Putative NECs (serotonergic cells associated with HNK1 epitope-immunoreactive neurites) are first visible at E18.5 in the medial gill epithelium. (e) Schematic (modified from [Tahara, 1988](#)) showing neural crest labelling by Dil injection at E5 (Piavis stages 11–12). (f–n) Two different embryos (f–j, k–n), showing Dil-labeled neural crest cells migrating ventrally (arrowheads, **g,l**) into the branchial arches, contributing to branchial arch basket and gill supporting cells (arrowheads, **j,n**), but not serotonergic cells (putative NECs) in the orobranchial epithelium (**i,j**) or gills (**n**). 5-HT, serotonin; Ba, branchial arch; Bab, branchial arch basket; En, endostyle; Gs, gill seam; Is, interbranchial septum; Nc, notochord; Nt, neural tube; Obc, orobranchial cavity. Scale-bar: 50 μ m.

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The following figure supplement is available for figure 4:

Figure supplement 1. Confirmation of successful targeting of pharyngeal arch-destined neural crest cells in all lamprey embryos analyzed for neural crest contribution to putative NECs.

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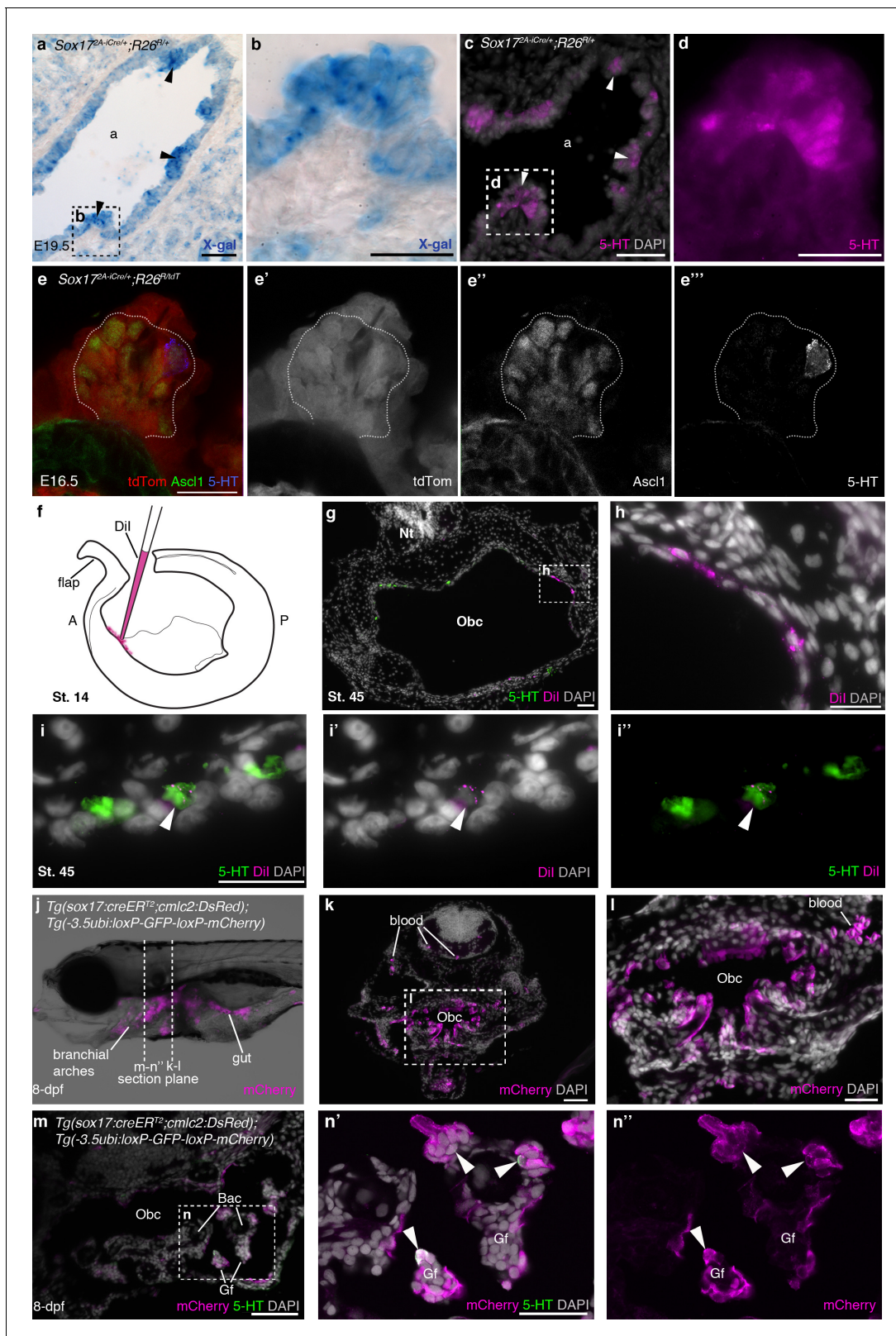


Figure 5. NECs are endoderm-derived, like PNECs. (a–d) In *Sox17^{2A-iCre/+};R26^{R/+}* mice, all endoderm-derived lineages, as well as vascular endothelial cells and the hematopoietic system, constitutively express β -galactosidase (Engert et al., 2009). Serial sections of an E19.5 *Sox17^{2A-iCre/+};R26^{R/+}* mouse lung show that X-gal labels PNECs (a,b; black arrowheads), whose identity is confirmed by serotonin expression (c,d; white arrowheads). The serotonin-positive cells are clearly all in the epithelium, which is entirely X-gal-positive, although there is some variation in staining level from cell to cell. (e–e''') A Figure 5 continued on next page

Figure 5 continued

high-power view of a cluster of Ascl1-expressing PNECs in a section of an E16.5 *Sox17^{2A-iCre/+};R26^{RtdTomato}* mouse lung, in which endoderm-derived lineages express tdTomato. Only the occasional PNEC is serotonin-positive at this stage. The Ascl1-expressing PNECs are tdTomato-positive, i.e., endoderm-derived. (f–i'') An endodermal contribution to putative NECs in *Xenopus* was investigated by performing focal Dil injections into the anterior endoderm at stage 14 (f), as described in **Chalmers and Slack (2000)**. At stage 45, Dil labels the endoderm lining the orobranchial cavity (g,h), and serotonergic cells (putative NECs, arrowheads) in the orobranchial epithelium (i–i''). (j–l) In *Tg(sox17:creER^{T2};cmlc2:DsRed);Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* zebrafish, the endoderm is labeled with mCherry and (m–n'') NECs in the gill filaments are mCherry-positive (arrowheads). 5-HT, serotonin; A, anterior; a, airway; Bac, branchial arch cartilage; Gf, gill filament; Obc, orobranchial cavity; P, posterior; tdTom, tdTomato. Scale-bars: 50 μ m in a,c,g,k,m; 25 μ m in b,d,h,i,l,n; 20 μ m in e.

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The following figure supplement is available for figure 5:

Figure supplement 1. The neural crest does not contribute to amniote PNECs.

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Cre expression and recombination in embryos from crosses between a *sox17:creER^{T2}* zebrafish driver line [*Tg(sox17:creER^{T2};cmlc2:DsRed)*; Joseph J. Lancman, Keith P. Gates, and P. Duc S. Dong, personal communication, March, 2017] and the switchable reporter line *Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* (**Mosimann et al., 2011**) (**Figure 5j–n''**). At 8-dpf, mCherry expression was seen in both gill NECs and putative NECs in the orobranchial epithelium: 147/331 serotonergic cells counted across six larvae (≥ 36 cells counted per fish) were mCherry-positive (**Figure 5n–n''**). [The relatively low labeling efficiency likely results from a lack of optimization of the 4-OHT dose for the *Tg(sox17:creER^{T2};cmlc2:DsRed)* driver in combination with this particular switchable reporter line (**Mosimann et al., 2011**).] These data demonstrate an endodermal origin in zebrafish for gill NECs, and also for putative NECs in the orobranchial epithelium. (Our genetic lineage-tracing data also confirm the endodermal origin of zebrafish gill filament epithelium, previously reported from vital dye fate-mapping experiments; **Warga and Nüsslein-Volhard, 1999**.)

Taken together, these results reveal the endodermal origin of gill NECs and putative NECs in the orobranchial epithelium. This supports the shared evolutionary ancestry of NECs with endoderm-derived PNECs, rather than neural crest-derived glomus cells.

NECs do not express Phox2b, which is essential for glomus cell development

It is formally possible that, despite their different embryonic origins, glomus cells and NECs could still be homologous cell types through activation of the same genetic network. Although the molecular basis of NEC development has not been investigated, the basic helix-loop-helix transcription factor Ascl1 (Mash1) is required for the formation of both PNECs (**Ito et al., 2000**) and glomus cells (**Kameda, 2005**). The homeodomain transcription factor Phox2b is also essential for glomus cell development (**Dauger et al., 2003**). However, we were unable to detect Phox2b-positive cells in embryonic zebrafish gills or orobranchial epithelium at 5-dpf (n = 4) or 7-dpf (n = 3), although Phox2b was expressed by a subset of cells in the hindbrain, as expected (**Coppola et al., 2012**) (**Figure 6a–f'**). Similarly, we could not detect any *Phox2*-positive cells in sea lamprey gills or orobranchial epithelia at E16 or E18 (n = 6), although *Phox2* was expressed in the epibranchial ganglia, and in patches of ectoderm and subjacent mesenchyme ventral to the epibranchial ganglia (**Figure 6g–j'**), in the same position as the hypobranchial placodes and associated ganglia identified in *Xenopus* (**Schlosser, 2003**).

We also found that PNECs lack Phox2b expression. In mouse embryos at E16.5, when PNECs can be identified by Ascl1 expression (serotonin is only expressed in a few PNECs at this stage), Phox2b was not seen in PNECs, despite expression in nearby intrinsic pulmonary ganglia (n = 2; **Figure 6k–l'**). Similarly, in chicken embryos at E12–E13.5, when scattered PNECs can be identified in the lung epithelium by serotonin immunoreactivity, Phox2b expression was not seen in the lung epithelium either by in situ hybridization or by immunostaining, although it could be detected in nearby intrinsic pulmonary ganglia (n = 3; **Figure 6m,m'**).

Overall, these data show that NECs (and PNECs) do not activate the same genetic network as glomus cells, since they lack expression of a transcription factor, Phox2b, which is essential for

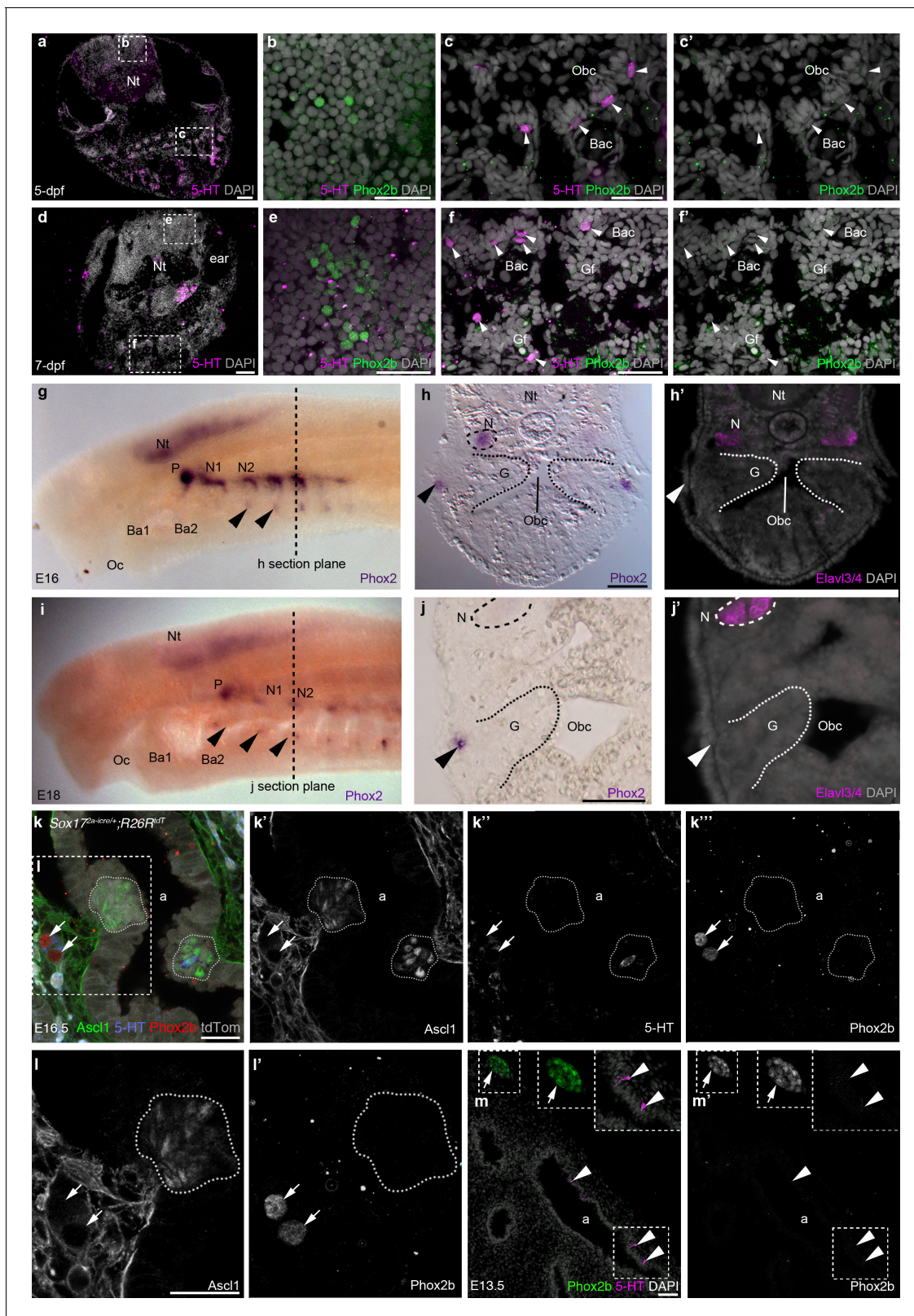


Figure 6. *Phox2b* expression is absent from gill and lung epithelia. (a–f') In wild-type zebrafish at 5- and 7-dpf, *Phox2b* is expressed by a subset of cells in the hindbrain (b,e), but not by gill NECs or putative NECs in the orobranchial epithelium (arrowheads; c–c',f–f'). (g–j') At E16 (g–h') and E18 (i–j') in the sea lamprey, *Phox2* expression is seen in the neural tube, the epibranchial (petrosal and nodose) ganglia (identified in section by the neuronal marker *Elavl3/4*), and in patches of ectoderm and subjacent mesenchyme ventral to the epibranchial ganglia (arrowheads). However, *Phox2* expression

Figure 6 continued on next page

Figure 6 continued

is absent from the gill epithelium, where putative NECs would be located. Dotted lines in panels **g** and **i** indicate the section plane in **h** and **j**. (**k–l'**) In a section of an E16.5 *Sox17^{2A-iCre/+};R26^{tdTomato}* mouse lung, *Phox2b* expression is seen in intrinsic pulmonary ganglia (arrows), but not in *Ascl1*/serotonin-positive PNECs located in the lung airway epithelium (dotted lines outline clusters of PNECs). (**m,m'**) In a section of an E13.5 chicken lung, *Phox2b* expression is seen in an intrinsic pulmonary ganglion (arrow), but not in serotonin-positive PNECs scattered in the lung airway epithelium (arrowheads). Insets show higher power views. a, airway; Ba, branchial arch; Bac, branchial arch cartilage; G, gill; Gf, gill filament; N, nodose ganglion; Nt, neural tube; Obc, orobranchial cavity; Oc, oral cavity; P, petrosal ganglion. Scale bars: 50 μ m in **a,d,h,j,m**; 25 μ m in **b,c,e,f,k,l**.

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glomus cell development (Dauger et al., 2003). Hence, NECs and glomus cells cannot be homologous cell types.

Candidate glomus cell homologues in fish: neural crest-derived chromaffin cells associated with pharyngeal arch blood vessels

Since NECs are endoderm-derived, we reasoned that neural crest-derived glomus cells must have evolved independently. Glomus cells are catecholaminergic, like the neural crest-derived chromaffin cells of the adrenal gland, which are also hypoxia-sensitive (reviewed by López-Barneo et al., 2016). Intriguingly, in lampreys, catecholaminergic (chromium salt-staining, i.e., 'chromaffin') cells were reported a century ago in association with large blood vessels not only in the trunk, but also as far rostrally as the second branchial arch, 'in the walls of the segmental veins as these run round the notochord' (Giacomini, 1902; Gaskell, 1912). This suggested to us the possibility that glomus cells could have evolved from catecholaminergic cells associated with blood vessels in anamniote pharyngeal arches, if such cells are neural crest-derived.

We first used immunostaining for the catecholaminergic marker tyrosine hydroxylase and the neurite-marker acetylated tubulin to confirm the existence of catecholaminergic cells, at least some of which may be innervated, in the walls of the anterior cardinal veins in the gill arches of ammocoete-stage sea lamprey (Figure 7a–c). It would be difficult to test whether the gill arch catecholaminergic cells are neural crest-derived: even if *Dil*-labeled embryos could be raised to ammocoete stages, the *Dil*-labeling would likely be very sparse. We reasoned that if present in lamprey, such cells might also be present in the zebrafish. We sectioned metamorphic juveniles and identified similar clusters of catecholaminergic cells in close association with blood vessels in the gill arches (Figure 7d–e''). To our knowledge, this is the first demonstration of the existence of catecholaminergic cells associated with gill arch blood vessels in a jawed anamniote. Importantly, these catecholaminergic cells were mCherry-positive, i.e., neural crest-derived, in *Tg(crestin:creER^{T2});Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* metamorphic juveniles (Figure 7e–e'') (51 such catecholaminergic cells were mCherry-positive across five juveniles [≥ 5 counted per fish]). This discovery suggests a new, speculative hypothesis for carotid body evolution, namely that it evolved in the amniote lineage via the aggregation of neural crest-derived catecholaminergic cells that were already associated with pharyngeal arch blood vessels in anamniotes.

Discussion

Gill NECs are endoderm-derived, suggesting shared ancestry with PNECs, not glomus cells

The evolutionary history of the hypoxia-sensitive cells that initiate amniote respiratory reflexes has been obscure, but is critical for our understanding of the transition from aquatic to terrestrial life. This involved a change from aquatic respiration in an environment with low oxygen solubility, to obligate air-breathing in an environment with more stable oxygen levels. It has been proposed that this transition was accompanied by a switch from a dispersed population of externally oriented hypoxia-sensitive cells in the gills that monitored the highly variable external oxygen levels, to one dominant site of hypoxia-sensitive cells that focused on monitoring internal oxygen states (Burlison and Milsom, 2003; Milsom and Burlison, 2007). Current hypotheses suggest that this change was accompanied by the evolution of the glomus cells of the carotid body from an ancestral population of gill NECs (e.g., Milsom and Burlison, 2007; Hempleman and Warburton, 2013; Jonz et al., 2016).

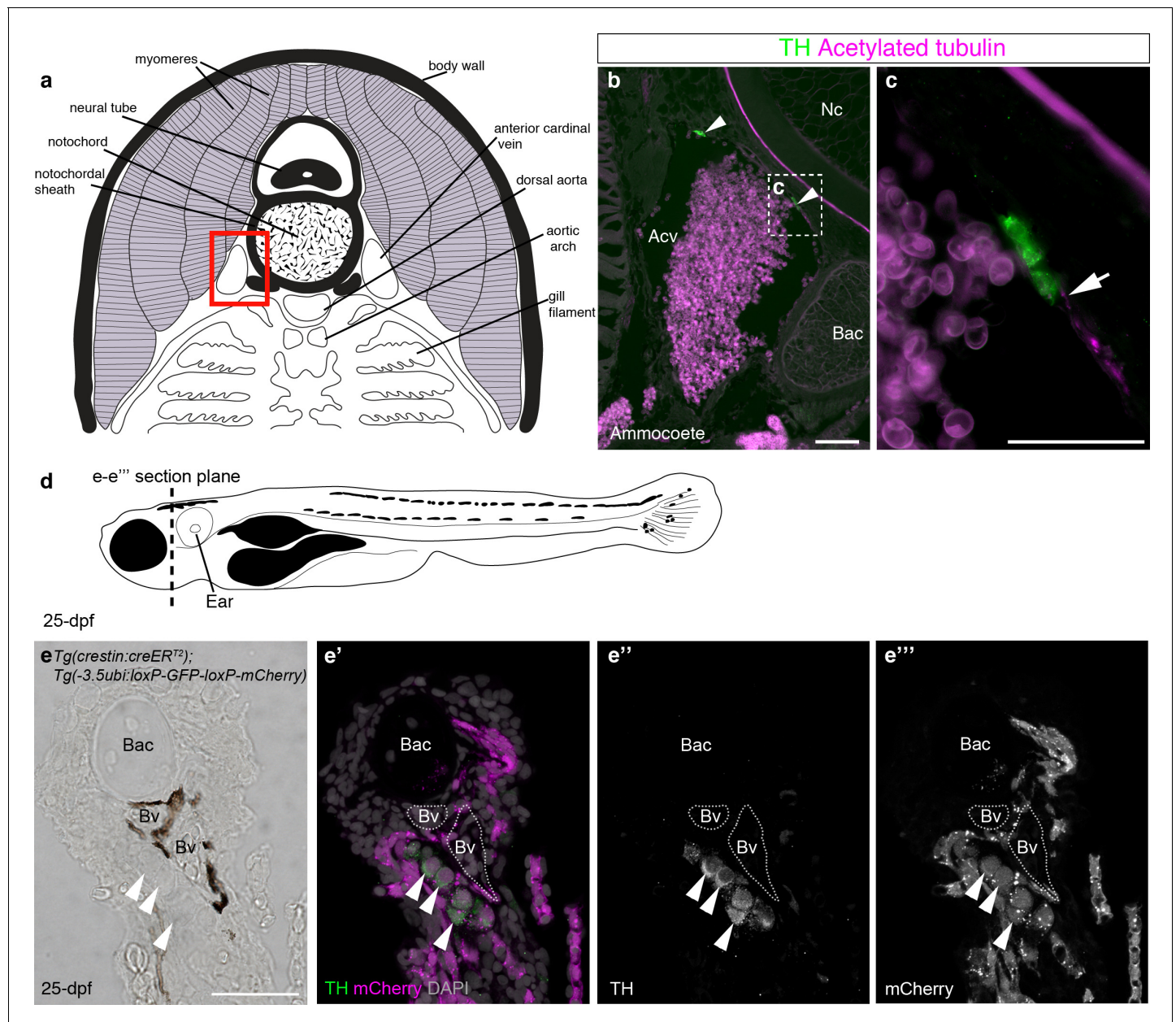


Figure 7. Catecholaminergic cells associated with gill arch blood vessels are neural crest-derived in zebrafish. (a) Schematic transverse section through ammocoete-stage lamprey gill arch (modified from [Ruppert et al., 2003](#)). Red box indicates region shown in b. (b,c) Tyrosine hydroxylase-positive (catecholaminergic) cells are present in the wall of the anterior cardinal vein (arrowheads), closely associated with acetylated tubulin-immunoreactive neurites (arrow). (d) Schematic 25-dpf zebrafish. Dotted line indicates transverse section plane through the gill basket in e-e'''. (e-e''') Tyrosine hydroxylase-positive (catecholaminergic) cells (arrowheads) seen adjacent to melanocyte-covered gill-filament blood vessels (dotted lines), are neural crest-derived (mCherry-positive) in 25-dpf *Tg(crestin:creER^{T2}); Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* zebrafish. Acv, anterior cardinal vein; Bac, branchial arch cartilage; Bv, blood vessel; Nc, notochord; TH, tyrosine hydroxylase. Scale-bars: 50 μ m in b; 25 μ m in c,e.

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This was entirely plausible, given their common association with pharyngeal arch arteries, afferent innervation by glossopharyngeal and/or vagal nerves, and hypoxia-sensitive K^+ currents ([López-Barneo et al., 1988](#); [Buckler, 1997](#); [Jonz et al., 2004](#); [Qin et al., 2010](#)). However, glomus cells are neural crest-derived ([Le Douarin et al., 1972](#); [Pearse et al., 1973](#); [Pardal et al., 2007](#)) and the hypothesis is not compatible with our demonstration that neural crest cells do not contribute to gill NECs in zebrafish, or their presumed homologues in *Xenopus* and lamprey, or to similar innervated

serotonergic cells (putative NECs) in the orobranchial epithelium of all three anamniote species. In contrast, we found that these cells are endoderm-derived, like PNECs, which differentiate in situ within pulmonary airway epithelia (Hoyt *et al.*, 1990; Rawlins *et al.*, 2009; Song *et al.*, 2012; Kuo and Krasnow, 2015) and whose endodermal origin we demonstrated using Sox17-Cre lineage-tracing in mouse. This suggests that gill NECs, and putative NECs in the orobranchial epithelium if these prove to be hypoxia-sensitive, more likely share a common ancestor with PNECs. Denervation experiments in various fishes (a shark, as well as some teleosts) have shown that hypoxia-sensitive and/or CO₂-sensitive chemoreceptors involved in ventilation responses are located in the orobranchial cavity, as well as in the gills, although their cellular identity has not been confirmed (reviewed by Milsom, 2012). Furthermore, putative NECs (identified by morphology, innervation and serotonin immunoreactivity, although not as yet shown to be hypoxia-sensitive) have been reported in the epithelium of the air-breathing organs (where present) of ray-finned fishes, lobe-finned lungfishes and amphibians (reviewed by Jonz *et al.*, 2016; Hsia *et al.*, 2013). Anamniote air-breathing organs likely evolved from out-pocketings of the caudal orobranchial epithelium after the evolution of gills and NECs (reviewed by Hsia *et al.*, 2013). Taken together, this leads us to speculate that hypoxia-sensitive NECs in the epithelia of the gills and orobranchial cavity of ancestral vertebrates were retained in the air-breathing organs of both anamniotes and amniotes. Testing this hypothesis for the evolutionary origin of PNECs must await evidence for the hypoxia-sensitivity of the putative NECs in the orobranchial epithelium of anamniotes.

Putative NECs have also been identified via serotonin immunoreactivity in the skin of developing zebrafish (Jonz and Nurse, 2006; Coccimiglio and Jonz, 2012) and of adult mangrove killifish, which respire through the skin as well as the gills (Regan *et al.*, 2011). Our genetic lineage-tracing data in zebrafish show that these cells are not neural crest-derived. If these serotonergic cells in the embryonic skin indeed prove to be hypoxia-sensitive NECs, then we suggest that NECs are likely a 'local epithelial' rather than uniquely endodermal cell type, i.e., that they can differentiate within epithelia of either ectodermal or endodermal origin, like taste buds (Barlow and Northcutt, 1995; Stone *et al.*, 1995) and the ameloblast (enamel-forming) layer of teeth (Soukup *et al.*, 2008).

It remained formally possible that, despite their different embryonic origins, gill NECs and glomus cells could be homologous cell types via activation of the same genetic network. However, we found that the transcription factor Phox2b, which is critical for glomus cell development (Dauger *et al.*, 2003), is not expressed by embryonic zebrafish gill NECs, putative NECs in the orobranchial epithelium, or their putative homologues in lamprey embryos. (We also found that mouse and chicken PNECs lack Phox2b expression.) Hence, NECs and glomus cells cannot be homologous cell types.

A new hypothesis for carotid body evolution

Since neural crest-derived glomus cells could not have evolved from endoderm-derived gill NECs, what is their evolutionary history? Glomus cells are catecholaminergic, and they are strikingly similar to the neural crest-derived chromaffin (i.e., catecholaminergic) cells of the adrenal medulla. For example, in fetal or neonatal mammals, adrenal chromaffin cells release catecholamines in direct response to hypoxia, like glomus cells (Comline and Silver, 1966; Cheung, 1989, 1990; Seidler and Slotkin, 1985; Thompson *et al.*, 1997). This 'non-neurogenic' response to hypoxia, in the absence of neural input, facilitates the transition to air-breathing in neonates by stimulating lung fluid absorption and regulating cardiovascular function (Seidler and Slotkin, 1985; Thompson *et al.*, 1997). The sensitivity of adrenal chromaffin cells to hypoxia is lost upon postnatal cholinergic innervation of the adrenal gland, although at least some hypoxia-responsive chromaffin cells persist in the adult adrenal medulla (García-Fernández *et al.*, 2007; Levitsky and López-Barneo, 2009) (also see López-Barneo *et al.*, 2016). Furthermore, hypoxia inhibits K⁺ currents in adrenal chromaffin cells, as in glomus cells (reviewed by López-Barneo *et al.*, 2016), and the 'set point' of hypoxia sensitivity is controlled in both glomus cells and adrenal chromaffin cells by mutual antagonism between the oxygen-regulated transcription factors hypoxia-inducible factor 1- α (Hif1 α /HIF1 α) and hypoxia-inducible factor 2- α (Hif2 α /HIF2 α) (Yuan *et al.*, 2013).

These similarities led us to re-visit century-old reports (Giacomini, 1902; Gaskell, 1912) of chromaffin (chromium salt-staining, i.e., catecholaminergic) cells associated with large branchial arch blood vessels in lamprey. We confirmed the existence of these catecholaminergic cells in ammocoete-stage sea lamprey, and went on to discover catecholaminergic cells associated with pharyngeal arch blood vessels in juvenile zebrafish, whose neural crest origin we demonstrated by genetic

lineage-tracing. We speculate that the carotid body may have evolved via the aggregation of such cells, and their subsequent acquisition of serotonergic properties and afferent innervation by glossopharyngeal and/or vagal afferents, such that they became incorporated into the afferent arm of respiratory reflexes. In order to test this hypothesis, it will be necessary to investigate whether these blood vessel-associated catecholaminergic cells in anamniotes secrete catecholamines in response to hypoxia. Glomus cells in the carotid body are enveloped by glial-like sustentacular cells, also neural crest-derived (Le Douarin *et al.*, 1972; Pearse *et al.*, 1973; Pardal *et al.*, 2007), which have been shown to act as adult stem cells for the production of new glomus cells under hypoxic conditions (Pardal *et al.*, 2007). Hypotheses for carotid body evolution must also take these cells into account.

The amphibian carotid labyrinth, a maze-like vascular expansion at the bifurcation of the carotid artery, is sensitive to oxygen levels (Ishii *et al.*, 1966) and considered a carotid body homologue (reviewed by Kusakabe, 2009). It contains both serotonergic and catecholaminergic cells, innervated by glossopharyngeal and/or vagal nerves (Reyes *et al.*, 2014), with both efferent and afferent synapses (reviewed by Kusakabe, 2009). Furthermore, the glomus cells in the carotid labyrinth are enveloped by fine processes of sustentacular cells (reviewed by Kusakabe, 2009). In adult amphibians, the carotid labyrinth is hypothesized to replace the NECs of the larval gills as the primary site of the chemosensors responsible for maintaining respiratory homeostasis (Kusakabe, 2002; Jonz and Nurse, 2006). Direct electrophysiological evidence is lacking for which cells are hypoxia-responsive, however, and their embryonic origin has not been established. Further investigation of the development and physiology of the carotid labyrinth in amphibians, and of the neural crest-derived catecholaminergic cells that we discovered in association with pharyngeal arch blood vessels in zebrafish, should help to test our new hypothesis for carotid body evolution.

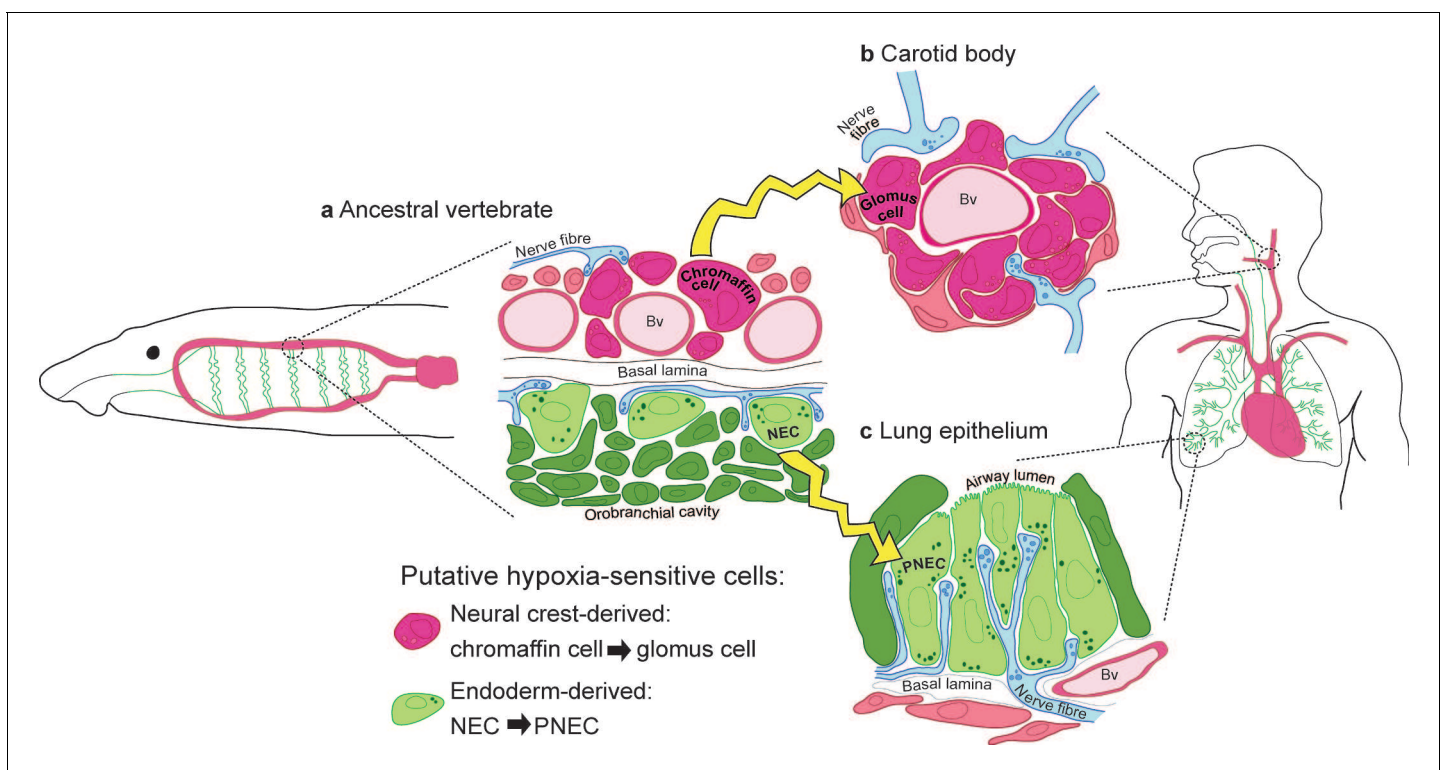


Figure 8. Model for the evolution of the hypoxia-sensitive cells involved in amniote respiratory reflexes. (a) Schematic ancestral vertebrate with internal gills. Neural crest-derived (magenta) chromaffin cells are associated with large blood vessels in the pharyngeal arches, while NECs differentiate within the endoderm-derived (green) epithelium lining the gills and orobranchial cavity. (b) During the transition to terrestrial life, the glomus cells of the carotid body evolved via the aggregation of neural crest-derived chromaffin cells (which must also have acquired serotonergic properties), while (c) NECs persisted as PNECs in lung airway epithelia. Yellow arrows indicate shared ancestry. Bv, blood vessel; NEC, neuroepithelial cell; PNEC, pulmonary neuroendocrine cell.

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A new model for amniote hypoxia-sensitive cell evolution

Given our lineage-tracing data, we present a new model for the evolution of the hypoxia-sensitive cells involved in amniote respiratory reflexes (**Figure 8**). We hypothesize that carotid body glomus cells evolved via the aggregation of neural crest-derived catecholaminergic (chromaffin) cells that were already associated with blood vessels in anamniote gill arches (and which must subsequently have acquired serotonergic properties and afferent innervation by glossopharyngeal and/or vagal nerves), while NECs differentiating in situ in the endoderm-derived epithelia of the gills and orobranchial cavity were retained as PNECs in lung airway epithelia. This model can be viewed as more parsimonious, since both embryonic lineage (neural crest versus endoderm) and function (physiological versus environmental oxygen monitoring) are maintained during the proposed evolutionary history of glomus cells and PNECs. Testing the model will require investigation of the physiology and hypoxia-responsiveness of anamniote gill arch blood vessel-associated catecholaminergic cells.

Materials and methods

Zebrafish lines

The following zebrafish (*Danio rerio*) lines were used: *Tg(-28.5sox10:cre);Tg(ef1a:loxP-DsRed-loxP-EGFP)* (Kague et al., 2012), *Tg(-4.9sox10:creER^{T2});Tg(βactin:loxP-SuperStop-loxP-DsRed)* (Mongera et al., 2013), *Tg(crestin:creER^{T2});Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* (Mosimann et al., 2011; Kaufman et al., 2016), *tfap2a^{mob};foxd3^{mos}* (Wang et al., 2011) and *Tg(sox17:creER^{T2};cmlc2:DsRed)* [created using the *sox17* promoter from Mizoguchi et al. (2008); Joseph J. Lancman, Keith P. Gates, and P. Duc S. Dong, personal communication, March, 2017]. Experiments using *Tg(-4.9sox10:creER^{T2});Tg(βactin:loxP-SuperStop-loxP-DsRed)* zebrafish were conducted in compliance with the regulations of the Regierungspräsidium Tübingen and the Max Planck Society. Experiments using all other zebrafish lines were conducted according to protocols approved by the Institutional Animal Care and Use Committees in facilities accredited by the Association for Assessment and Accreditation of Laboratory Animal Care International (AAALAC). All zebrafish were fixed overnight at 4°C in 4% paraformaldehyde in phosphate-buffered saline (PBS), except for *Tg(sox17:creER^{T2};cmlc2:DsRed);Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* zebrafish, which were fixed overnight at 4°C in 4% paraformaldehyde in 0.1 M Pipes, 1 mM MgSO₄, 2 mM EGTA, pH 7).

To induce Cre activity and recombination in *Tg(-4.9sox10:creER^{T2});Tg(βactin:loxP-SuperStop-loxP-DsRed)* zebrafish, embryos were dechorionated at 16 hr post-fertilization (hpf) and treated with 5 μM 4-hydroxytamoxifen (4-OHT; Sigma-Aldrich, St. Louis, MO) for 8 hr. As reported in Mongera et al. (2013), 4-OHT treatment of this line for 8 hr from 16 hpf is very effective in yielding Cre-induced recombination in the branchial arches, and was used in Mongera et al. (2013) to demonstrate the neural crest origin of gill pillar cells. To induce Cre activity and recombination in *Tg(crestin:creER^{T2});Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* zebrafish, embryos were treated with 20 μM 4-OHT in ethanol at 50% epiboly and again at 24 hpf. To induce Cre activity and recombination in *Tg(sox17:creER^{T2};cmlc2:DsRed);Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* zebrafish, embryos were treated for 3 hr from 5 hpf with 10 μM 4-OHT.

Xenopus neural fold grafts

Experiments using *Xenopus laevis* were conducted in accordance with the UK Animals (Scientific Procedures) Act 1986, with appropriate personal and project licences in place where necessary. Embryos were obtained by in vitro fertilization and initially kept at 14°C in 0.1% modified Barth's saline (MBS). For grafted embryos that would be grown to stage 53 (after the onset of independent feeding), CMV-GFP transgenic embryos (Marsh-Armstrong et al., 1999) were used as donors and the embryos were grafted and reared at the European *Xenopus* Resource Centre (University of Portsmouth, UK). For grafted embryos that were to be grown to embryonic stages 41–45, GFP-positive donor embryos were made by injecting cyto-GFP mRNA into one cell at the two-cell stage, or two cells at the four-cell stage. Briefly, embryos were de-jellied in 2% cysteine and washed several times in 0.1% MBS before being transferred and positioned for injection in a mesh-lined Petri dish filled with 4% Ficoll. Injected embryos were allowed to recover in 4% Ficoll for at least 1 hr before being transferred to 0.1% MBS.

De-jellied GFP-positive embryos and wild-type embryos were allowed to grow to stage 13–17 at 14–18°C. For grafting, embryos were moved to 18 mm Petri dishes lined with plasticine or 1% agarose with depressions and containing a high-salt transplantation solution (1x MBSH: 1x MBS, 0.7 mM CaCl_2 , 0.02 mM NaCl, supplemented with 2 mM CaCl_2 and 2.5 mg/ml gentamycin [Sigma-Aldrich]). The region of the neural folds containing premigratory branchial and vagal neural crest (Sadaghiani and Thiébaud, 1987) was removed unilaterally from stage 13–15 wild-type hosts and replaced with GFP-positive tissue from the same region of donor embryos (Figure 3d). The grafted tissue was held in place with a small piece of glass coverslip while embryos recovered in transplantation solution for at least 2 hr, before being moved to 0.1% MBS and reared at 18°C. Embryos were overdosed in MS222 (Sigma-Aldrich) in PBS before being fixed in 4% paraformaldehyde in PBS overnight at 4°C.

Lamprey Dil injections

Experiments using sea lamprey (*Petromyzon marinus*) were conducted according to protocols approved by the California Institute of Technology Institutional Animal Care and Use Committee. Eggs were collected from adults and fertilized as described (Nikitina et al., 2009). Embryos were maintained at 18°C in 0.1x or 1x Marc's modified Ringer's (MMR) solution. Dil labeling was performed as described (Nikitina et al., 2009) with some modifications. Briefly, E5 embryos (Piavis stages 11–12) were manually dechorionated in 0.1x MMR, then immobilized and oriented in 18-mm Petri dishes that were lined with a fine mesh. Embryos were pressure-injected into the dorsal neural tube using glass capillary tubes filled with 0.5 mg/ml of Cell Tracker-CM-Dil (Invitrogen, Carlsbad, CA) diluted in 0.3 M sucrose (from a 5 mg/ml stock diluted in ethanol). They were allowed to recover for 24 hr, then individually transferred to an uncoated Petri dish containing 1x MMR and allowed to develop to E19 (Piavis stage 17). Embryos were periodically checked and imaged throughout, then fixed in 4% paraformaldehyde in PBS for 1 hr at room temperature.

Transgenic mouse lines

The following transgenic mouse lines were used: *Wnt1-cre;R26R-YFP* (Danielian et al., 1998; Srinivas et al., 2001), *Sox17^{2A-iCre};R26^{R/+}* (Engert et al., 2009; Soriano, 1999) and *Sox17^{2A-iCre};R26R^{tdTomato}* (Engert et al., 2009; Madisen et al., 2010). Experiments using these mice were conducted in accordance with the UK Animals (Scientific Procedures) Act 1986, with appropriate personal and project licences in place. Embryos were dissected at appropriate stages and fixed at 4°C overnight in 4% paraformaldehyde in PBS.

Chicken neural tube grafts

Experiments using chicken (*Gallus gallus domesticus*) embryos were conducted in accordance with the UK Animals (Scientific Procedures) Act 1986, with appropriate personal and project licences in place where necessary. Fertilized wild-type chicken eggs were obtained from Henry Stewart and Co. Ltd., Norfolk, UK. Fertilized GFP-transgenic chicken eggs (McGrew et al., 2008) were obtained from the Roslin Institute Transgenic Chicken Facility (Edinburgh, UK), which is funded by the Wellcome Trust and the BBSRC. Fertilized wild-type and GFP-transgenic eggs were incubated in a humidified atmosphere at 38°C for approximately 1.5 days to reach 8–11 somites. The neural tube and associated neural folds between the level of somite one and the caudal end of the seventh somite were dissected from a wild-type host and replaced with the equivalent tissue from a GFP-transgenic donor embryo. At E14, embryos were decapitated and the lungs dissected out and fixed overnight in 4% paraformaldehyde in PBS. At E16.5, embryos were decapitated and fixed overnight in 4% paraformaldehyde in PBS; the lungs were dissected after fixation. The lungs were dehydrated through an ethanol series into 100% ethanol for storage.

Xenopus Dil injections

Stage 14 *Xenopus laevis* embryos were fixed in place in a plasticine dish filled with 1x MBSH supplemented with 2 mM CaCl_2 , and the endoderm was exposed by cutting a flap into the anterior neural plate (leaving it attached on the anterior side) with tungsten needles, which was folded back to expose the endoderm. A stock solution of 2 mg/ml Cell Tracker-CM-Dil (Invitrogen) in ethanol was diluted 1:10 in 10% sucrose and microinjected into the anterior endoderm (regions 1 and 5 of

Chalmers and Slack, 2000) using a glass electrode whose tip was approximately 20 μm in diameter. The endoderm of each embryo was injected at three to five sites. The flap of the neural plate was then folded back in place and pressed down with a small piece of glass coverslip supported on plasticine feet until it healed back in place (approximately 1–2 hr). Embryos were then transferred into 0.1 x MBS containing 25 mg/l gentamicin (Sigma-Aldrich), 400 mg/l penicillin (Sigma-Aldrich) and 400 mg/l streptomycin sulfate (Sigma-Aldrich). At stage 45, tadpoles were overdosed in MS222 (Sigma-Aldrich) in PBS before being fixed in 4% paraformaldehyde in PBS at 4°C overnight for up to several days, then transferred to PBS.

Ammocoete lamprey

An ammocoete lamprey was euthanized by MS222 (Sigma-Aldrich) overdose, fixed in modified Carnoy's solution (six volumes ethanol: three volumes 37% formaldehyde: 1 volume glacial acetic acid) and dehydrated through an ethanol series into 100% ethanol.

Embedding and immunostaining

Dil-labeled lamprey embryos, grafted *Xenopus* embryos, some *Tg(-28.5sox10:cre;ef1a:loxP-DsRed-loxP-EGFP)*, *tfap2a^{mob};foxd3^{mos}* zebrafish embryos and their wild-type siblings were dehydrated from PBS into 100% methanol and transferred to 100% isopropanol overnight at 4°C. Embryos were transferred to 1:1 isopropanol:chloroform for 1 hr at 4°C and then to 100% chloroform for 2 hr at –20°C. After warming to room temperature, embryos were transferred to 1:1 chloroform:paraffin wax (Raymond A. Lamb Ltd., Thermo Fisher Scientific, Waltham, MA) at 60°C for 30 min, followed by three 30-min incubations and an overnight incubation at 60°C in paraffin wax. Embryos were embedded in plastic molds and sectioned at 6 μm using a rotary microtome.

Dil-labeled *Xenopus* embryos, *Tg(crestin:creER^{T2});Tg(-3.5ubi:loxP-GFP-loxP-mCherry)*, *Tg(-4.9sox10:creER^{T2});Tg(β actin:loxP-SuperStop-loxP-DsRed)*, some *Tg(-28.5sox10:cre);Tg(ef1a:loxP-DsRed-loxP-EGFP)* and *Tg(sox17:creER^{T2};cmlc2:DsRed);Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* zebrafish in PBS were sucrose-protected before being embedded in 7.5–20% gelatin in plastic molds, flash-frozen in liquid nitrogen and cryosectioned at 6 μm .

Mouse embryos and wild-type zebrafish embryos were sucrose-protected before being embedded in O.C.T. compound (Tissue-Tek, Sakura Finetek, Torrance, CA) in plastic molds, flash-frozen in isopentane on dry ice and cryosectioned at 10–15 μm .

Grafted chicken lungs and the ammocoete lamprey in 100% ethanol were cleared in HistoSol (National Diagnostics, Atlanta, GA) and incubated in 1:1 HistoSol: paraffin wax (Raymond A. Lamb Ltd.) for 30 min at 60°C, followed by three 30-min incubations and an overnight incubation in paraffin wax at 60°C. They were then embedded in plastic molds and sectioned at 6–10 μm using a rotary microtome.

For immunostaining on paraffin wax sections, slides were de-waxed in HistoSol and rehydrated into PBS through a graded ethanol series. Cryosections were allowed to warm to room temperature and washed in PBS. When necessary, gelatin was removed by dipping slides in PBS warmed to 37°C. All anti-serotonin antibodies used required antigen retrieval, which was performed by heating the slides for 30 s in a microwave in 10 mM sodium citrate buffer solution (pH 6), followed by two washes in PBS. Immunostaining was performed as described (**Nikitina et al., 2009**) with slight modifications: slides were incubated overnight at 4°C or at room temperature in primary antibody in blocking solution (10% sheep, goat or donkey serum, as appropriate, in PBS with 0.1% Triton X-100); secondary antibodies were incubated at room temperature for 2 hr or overnight at 4°C. For horse-radish peroxidase detection, slides were incubated in 0.3 mg/ml diaminobenzidine, 0.02% H₂O₂, 0.05% Triton X-100 in PBS. After immunostaining, sections were counterstained with the nuclear marker DAPI (1 ng/ml) (Invitrogen) and mounted in Fluoromount G (Southern Biotech, Birmingham, AL).

For whole-mount immunostaining, *Tg(crestin:creER^{T2});Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* zebrafish embryos were incubated for 2 hr in blocking buffer (PBS with 4% bovine serum albumin, 0.3% Triton X-100, 0.02% sodium azide) prior to overnight incubation at 4°C with primary antibodies diluted in blocking buffer. Embryos were washed for 2 hr at room temperature in PBS with 0.3% Triton X-100, then incubated overnight at 4°C in blocking buffer containing secondary antibodies

diluted 1:200 and 1 mg/ml DAPI (Invitrogen) diluted 1:200. After washing for 2 hr in PBS with 0.3% Triton X-100, embryos were suspended in 80% glycerol before mounting.

Antibodies

Primary antibodies were used against the following antigens: acetylated tubulin [1:250 mouse IgG2b, clone 6-11-B1, T7451 Sigma-Aldrich; previously used in the sea lamprey, e.g., *Barreiro-Iglesias et al. (2008a)*], Ascl1 (Mash1) [1:200 mouse IgG1 (*Lo et al., 1991*), kind gift of F. Guillemot, NIMR, London, UK; 1:100 mouse IgG1, #556604 BD Biosciences, San Jose, CA], DsRed2 (1:100 mouse IgG1, sc-101526 Santa Cruz Biotechnology, Dallas, TX), Elavl3/4 (HuC/D) (1:500 mouse IgG2b, A-21271 Invitrogen), GFP (1:500 rabbit, A-6455 Invitrogen; 1:500 mouse IgG1, #1814460001 Roche, Basel, Switzerland; 1:250 goat, ab6662 Abcam [Cambridge, UK]; 1:150 chicken, ab13970 Abcam), HNK-1 carbohydrate epitope (*Abo and Balch, 1981; Voshol et al., 1996*) [for zebrafish neurites (*Metcalfe et al., 1990*): 1:100 mouse IgG1, ZN-12 Developmental Studies Hybridoma Bank; for lamprey neurites (*Barreiro-Iglesias et al., 2008b*): 1:50 mouse IgM, 3H5 Developmental Studies Hybridoma Bank], mCherry (1:250 mouse IgG1, #632543 Clontech Takara Bio USA Inc., Mountain View, CA; 1:200 goat, orb11618 Biorbyt, Cambridge, UK), serotonin (5-hydroxytryptamine, 5-HT) [1:100 (whole-mount) or 1:250 (sections) rabbit, S5545 Sigma-Aldrich, previously used in zebrafish, e.g., *Kuscha et al. (2012)*, bullfrog (*Reyes et al., 2014*) and Arctic lamprey (*Suzuki et al., 2015*); 1:100 rat, MAB352 Merck Millipore, Temecula, CA, previously used in zebrafish (*Sundvik et al., 2013*); 1:250 goat, ab66047 Abcam], Phox2b [1:500 rabbit, kind gift of Jean-François Brunet, Institut de Biologie de l'École Normale Supérieure, Paris, France; previously used in zebrafish (*Coppola et al., 2012*); Tubb3 (neuronal β -III tubulin) (1:500 mouse IgG2a, clone TUJ1, MMS-435P Covance BioLegend, San Diego, CA), and tyrosine hydroxylase [1:250, rabbit, AB152 Merck Millipore; previously used in zebrafish, e.g., *Yamamoto et al. (2011)*, and sea lamprey, e.g., *Barreiro-Iglesias et al. (2008a)*]. (The Developmental Studies Hybridoma Bank was developed under the auspices of the NICHD and is maintained by the University of Iowa, Department of Biological Sciences, Iowa City.) Appropriately matched AlexaFluor or horse-radish peroxidase-conjugated secondary antibodies were obtained from Molecular Probes/Invitrogen.

Whole-mount in situ hybridization

The lamprey (*P. marinus*) *Phox2* clone (*Häming et al., 2011*) was a kind gift of Marianne Bronner (Caltech, Pasadena, CA, USA). Whole-mount in situ hybridization on lamprey embryos was performed as described (*Nikitina et al., 2009*). After whole-mount in situ hybridization, embryos were incubated in PBS with 5% sucrose for 4 hr at room temperature, followed by incubation overnight at 4°C in 15% sucrose in PBS. Embryos were transferred into pre-warmed 7.5% gelatin in 15% sucrose in PBS and incubated for 1–4 hr at 37°C, then oriented and embedded in molds, frozen by immersion in a dry ice-isopentane solution for 30 s, and cryosectioned at 12–16 μ m. Gelatin was removed from the slides by a 5-min incubation in PBS pre-warmed to 37°C.

In situ hybridization on paraffin wax sections

The chicken *Phox2b* clone (*Stanke et al., 1999*) was a kind gift of Jean-François Brunet (Institut de Biologie de l'École Normale Supérieure, Paris, France). For in situ hybridization on paraffin wax sections, slides were de-waxed in Histosol (National Diagnostics) and rehydrated into diethylpyrocarbonate (DEPC)-treated (Sigma-Aldrich) PBS through a graded ethanol series. In situ hybridization was performed on sections as described (*Miller et al., 2017*).

X-gal staining

For X-gal staining on cryosections of mouse tissue, the following staining solution was added to slides prior to incubation at 37°C for 2 hr: 5 mM $K_3Fe(CN)_6$, 5 mM $K_4Fe(CN)_6$, 2.7 mM $MgCl_2$ in PBS, supplemented with 75 mg/ml X-gal in dimethyl sulfoxide (DMSO).

Alcian blue plus hematoxylin and eosin staining

For Alcian blue plus hematoxylin and eosin staining on paraffin sections, slides were de-waxed in Histosol and rehydrated into water through a graded ethanol series. Slides were rinsed in 3% acetic acid before staining in 2 mg/ml Alcian blue (Searle Diagnostic, High Wycombe, UK) in 3% acetic acid

for at least 30 min. Slides were then rinsed in water and treated with 0.3% NaHCO₃, followed by another rinse in running water and staining in Mayer's hematoxylin (Sigma-Aldrich) for 10 min. Slides were stained in 1% aqueous eosin Y solution (BDH) for 8 min, then washed again in running water before dehydration through an ethanol series into 100% ethanol. After washing in HistoSol, slides were mounted with DPX (BDH).

Image capture and processing

Whole-mount images were taken using a Leica MZFLIII microscope (Leica Microsystems, Wetzlar, Germany) fitted with a QImaging MicroPublisher 5.0 RTV camera and QCapture Pro 6.0 software (QImaging, Surrey, BC, Canada); a Zeiss AxioSkop2 microscope fitted with a Zeiss AxioCam HRc camera and Zeiss AxioVision Rel. 4.8 software (Carl Zeiss, Oberkochen, Germany); an Olympus MVX10 microscope (Olympus Corporation, Tokyo, Japan) fitted with a Zeiss AxioCam HRc camera and Zeiss AxioVision Rel. 4.8 software; an Olympus 1 × 71 inverted microscope fitted with a Hamamatsu ORCA-R2 monochrome camera and HCLImage software (Hamamatsu Photonics, Hamamatsu, Japan); a Zeiss LSM 710 confocal microscope with Zeiss ZEN software; and a Zeiss 710 confocal microscope with Zeiss LSM Image Browser (version 4.2.0.121) software, which was used to create three-dimensional images and stack movies. Images of sections were taken using a Zeiss AxioSkop 2 MOT microscope fitted with a QImaging Retiga 2000R camera, a Qimaging RGB pancake and QCapture Pro 6.0 software; a Zeiss Scope.A1 microscope fitted with a Zeiss AxioCam MRm camera and Zeiss ZEN 2012 (blue edition) software; and a Zeiss LSM 780 confocal microscope with Zeiss ZEN 2011 (black edition) software. All images were further processed in Photoshop CS4 (Adobe Systems Inc., San Jose, CA) and/or ImageJ 1.50i software (NIH, Bethesda, MD).

Statistical analysis

Data analysis and statistical tests were performed using Microsoft Excel and GraphPad Prism 7 software (GraphPad Software, Inc., La Jolla, CA). Data sets were tested for normality using the Shapiro-Wilk test ($\alpha = 0.05$) and for equality of variance using an F test ($p=0.38$), and compared using an unpaired two-tailed Student's t-test. Data are presented as mean \pm standard deviation (s.d.).

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Author contributions

DH, conceived and designed the experiments with CVHB, performed all experiments except the *Xenopus* Dil injections reported in Figure 5e-i' (GS), the chicken neural fold grafts reported in Figure 5-figure supplement 1c-e' (AJB), the whole-mount zebrafish immunostaining reported in Figure 5-figure supplement 2 (KPG) and the chicken Phox2b immunostaining reported in Figure 6m,m' (BJ), analyzed all the results except the zebrafish skin NEC data (KPG), prepared all the figures and wrote the manuscript with CVHB; AJB, performed the chicken neural fold grafts reported in Figure 5-figure supplement 1c-e', provided *Wnt1:cre;R26R:YFP* mouse embryos and commented on the manuscript; GS, performed the *Xenopus* Dil injections reported in Figure 5e-i' and commented on the manuscript; KPG, performed, imaged and analyzed the whole-mount zebrafish immunostaining data reported in Figure 5-figure supplement 2, provided (with JLL and PDSD) 4-OHT-treated *Tg(sox17:creER^{T2};cmlc2:DsRed);Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* zebrafish larvae, and commented on the

manuscript; BJ, performed and imaged the chicken Phox2b immunostaining reported in Figure 6m, m', and commented on the manuscript; AM, provided 4-OHT-treated *Tg(-4.9sox10:creER^{T2});Tg(βactin:loxP-SuperStop-loxP-DsRed)* zebrafish larvae and commented on the manuscript; SF, provided *Tg(-28.5sox10:cre);Tg(ef1a:loxP-DsRed-loxP-EGFP)* zebrafish larvae and commented on the manuscript; GU, provided (with EWK) *tfap2a^{mob};foxd3^{mos}* zebrafish larvae and commented on the manuscript; EWK, provided (with GU) *tfap2a^{mob};foxd3^{mos}* zebrafish larvae and commented on the manuscript; CKK, provided (with CM and LIZ) 4-OHT-treated *Tg(crestin:creER^{T2});Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* zebrafish larvae and juveniles, and commented on the manuscript; CM, provided (with CKK and LIZ) 4-OHT-treated *Tg(crestin:creER^{T2});Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* zebrafish larvae and juveniles, and commented on the manuscript; LIZ, provided (with CKK and CM) 4-OHT-treated *Tg(crestin:creER^{T2});Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* zebrafish larvae and juveniles, and commented on the manuscript; JKL, provided (with KPG and PDSD) 4-OHT-treated *Tg(sox17:creER^{T2};cmlc2:DsRed);Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* zebrafish larvae, and commented on the manuscript; PDSD, provided (with KPG and JKL) 4-OHT-treated *Tg(sox17:creER^{T2};cmlc2:DsRed);Tg(-3.5ubi:loxP-GFP-loxP-mCherry)* zebrafish larvae, and commented on the manuscript; HL, provided access to the *Sox17^{2A-iCre}* mouse line and commented on the manuscript; AST, provided *Sox17^{2A-iCre/+};R26^{R/+}* mouse embryos and cryosections of *Sox17^{2A-iCre/+};R26^RtdTomato* mouse embryos, and commented on the manuscript; CVHB, conceived and designed the experiments with DH and wrote the manuscript with DH

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Ethics

Animal experimentation: Experiments using *Tg(-4.9sox10:creER^{T2});Tg(βactin:loxP-SuperStop-loxP-DsRed)* zebrafish were conducted in compliance with the regulations of the Regierungspräsidium Tübingen and the Max Planck Society. Experiments using all other zebrafish lines were conducted according to protocols approved by the Institutional Animal Care and Use Committees in facilities accredited by the Association for Assessment and Accreditation of Laboratory Animal Care International (AAALAC). Experiments using *Xenopus laevis* were conducted in accordance with the UK Animals (Scientific Procedures) Act 1986, with appropriate personal and project licences in place where necessary. Experiments using sea lamprey (*Petromyzon marinus*) were conducted according to protocols approved by the California Institute of Technology Institutional Animal Care and Use Committee. Experiments using transgenic mice were conducted in accordance with the UK Animals (Scientific Procedures) Act 1986, with appropriate personal and project licences in place. Experiments using chicken (*Gallus gallus domesticus*) embryos were conducted in accordance with the UK Animals (Scientific Procedures) Act 1986, with appropriate personal and project licences in place where necessary.

References

- Abo T, Balch CM. 1981. A differentiation antigen of human NK and K cells identified by a monoclonal antibody (HNK-1). *Journal of Immunology* **127**:1024–1029. PMID: 6790607
- Akhtar S, Schonthaler HB, Bron AJ, Dahm R. 2008. Formation of stromal collagen fibrils and proteoglycans in the developing zebrafish cornea. *Acta Ophthalmologica* **86**:655–665. doi: 10.1111/j.1600-0420.2007.01135.x, PMID: 18221494
- Barlow LA, Northcutt RG. 1995. Embryonic origin of amphibian taste buds. *Developmental Biology* **169**:273–285. doi: 10.1006/dbio.1995.1143, PMID: 7750643
- Barrallo-Gimeno A, Holzschuh J, Driever W, Knapik EW. 2004. Neural crest survival and differentiation in zebrafish depends on *mont blanc/tfap2a* gene function. *Development* **131**:1463–1477. doi: 10.1242/dev.01033, PMID: 14985255
- Barreiro-Iglesias A, Aldegunde M, Anadón R, Rodicio MC. 2009. Extensive presence of serotonergic cells and fibers in the peripheral nervous system of lampreys. *The Journal of Comparative Neurology* **512**:478–499. doi: 10.1002/cne.21914, PMID: 19025991
- Barreiro-Iglesias A, Villar-Cerviño V, Villar-Cheda B, Anadón R, Rodicio MC. 2008a. Neurochemical characterization of sea lamprey taste buds and afferent gustatory fibers: presence of serotonin, calretinin, and CGRP immunoreactivity in taste bud bi-ciliated cells of the earliest vertebrates. *The Journal of Comparative Neurology* **511**:438–453. doi: 10.1002/cne.21844, PMID: 18831528

- Barreiro-Iglesias A**, Villar-Cheda B, Abalo XM, Anadón R, Rodicio MC. 2008b. The early scaffold of axon tracts in the brain of a primitive vertebrate, the sea lamprey. *Brain Research Bulletin* **75**:42–52. doi: [10.1016/j.brainresbull.2007.07.020](https://doi.org/10.1016/j.brainresbull.2007.07.020), PMID: [18158094](https://pubmed.ncbi.nlm.nih.gov/18158094/)
- Branchfield K**, Nantie L, Verheyden JM, Sui P, Wienhold MD, Sun X. 2016. Pulmonary neuroendocrine cells function as airway sensors to control lung immune response. *Science* **351**:707–710. doi: [10.1126/science.aad7969](https://doi.org/10.1126/science.aad7969), PMID: [26743624](https://pubmed.ncbi.nlm.nih.gov/26743624/)
- Buckler KJ**. 1997. A novel oxygen-sensitive potassium current in rat carotid body type I cells. *The Journal of Physiology* **498**:649–662. doi: [10.1113/jphysiol.1997.sp021890](https://doi.org/10.1113/jphysiol.1997.sp021890), PMID: [9051577](https://pubmed.ncbi.nlm.nih.gov/9051577/)
- Burleson ML**, Mercer SE, Wilk-Blaszczak MA. 2006. Isolation and characterization of putative O₂ chemoreceptor cells from the gills of channel catfish (*Ictalurus punctatus*). *Brain Research* **1092**:100–107. doi: [10.1016/j.brainres.2006.03.085](https://doi.org/10.1016/j.brainres.2006.03.085), PMID: [16690040](https://pubmed.ncbi.nlm.nih.gov/16690040/)
- Burleson ML**, Milsom WK. 2003. Comparative aspects of O₂ chemoreception. Anatomy, physiology, and environmental applications. In: Lahiri S, Semenza GL, Prabhakar NR (Eds). *Oxygen Sensing: Responses and Adaptation to Hypoxia*. New York: Marcel Dekker, Inc. p. 685–707.
- Burns AJ**, Delalande J-M. 2005. Neural crest cell origin for intrinsic ganglia of the developing chicken lung. *Developmental Biology* **277**:63–79. doi: [10.1016/j.ydbio.2004.09.006](https://doi.org/10.1016/j.ydbio.2004.09.006), PMID: [15572140](https://pubmed.ncbi.nlm.nih.gov/15572140/)
- Chalmers AD**, Slack JM. 2000. The *Xenopus* tadpole gut: fate maps and morphogenetic movements. *Development* **127**:381–392. PMID: [10603354](https://pubmed.ncbi.nlm.nih.gov/10603354/)
- Cheung CY**. 1989. Direct adrenal medullary catecholamine response to hypoxia in fetal sheep. *Journal of Neurochemistry* **52**:148–153. doi: [10.1111/j.1471-4159.1989.tb10909.x](https://doi.org/10.1111/j.1471-4159.1989.tb10909.x), PMID: [2908885](https://pubmed.ncbi.nlm.nih.gov/2908885/)
- Cheung CY**. 1990. Fetal adrenal medulla catecholamine response to hypoxia-direct and neural components. *The American Journal of Physiology* **258**:R1340–R1346. PMID: [1972870](https://pubmed.ncbi.nlm.nih.gov/1972870/)
- Coccimiglio ML**, Jonz MG. 2012. Serotonergic neuroepithelial cells of the skin in developing zebrafish: morphology, innervation and oxygen-sensitive properties. *Journal of Experimental Biology* **215**:3881–3894. doi: [10.1242/jeb.074575](https://doi.org/10.1242/jeb.074575), PMID: [22855620](https://pubmed.ncbi.nlm.nih.gov/22855620/)
- Comline RS**, Silver M. 1966. The development of the adrenal medulla of the foetal and new-born calf. *The Journal of Physiology* **183**:305–340. doi: [10.1113/jphysiol.1966.sp007868](https://doi.org/10.1113/jphysiol.1966.sp007868), PMID: [5949627](https://pubmed.ncbi.nlm.nih.gov/5949627/)
- Coppola E**, D'autréaux F, Nomaksteinsky M, Brunet J-F. 2012. *Phox2b* expression in the taste centers of fish. *The Journal of Comparative Neurology* **520**:3633–3649. doi: [10.1002/cne.23117](https://doi.org/10.1002/cne.23117), PMID: [22473338](https://pubmed.ncbi.nlm.nih.gov/22473338/)
- Cutz E**, Pan J, Yeger H, Domnik NJ, Fisher JT. 2013. Recent advances and controversies on the role of pulmonary neuroepithelial bodies as airway sensors. *Seminars in Cell & Developmental Biology* **24**:40–50. doi: [10.1016/j.semcdb.2012.09.003](https://doi.org/10.1016/j.semcdb.2012.09.003), PMID: [23022441](https://pubmed.ncbi.nlm.nih.gov/23022441/)
- Danielian PS**, Muccino D, Rowitch DH, Michael SK, McMahon AP. 1998. Modification of gene activity in mouse embryos *in utero* by a tamoxifen-inducible form of cre recombinase. *Current Biology* **8**:1323–S2. doi: [10.1016/S0960-9822\(07\)00562-3](https://doi.org/10.1016/S0960-9822(07)00562-3), PMID: [9843687](https://pubmed.ncbi.nlm.nih.gov/9843687/)
- Dauger S**, Pattyn A, Lofaso F, Gaultier C, Goridis C, Gallego J, Brunet J-F. 2003. *Phox2b* controls the development of peripheral chemoreceptors and afferent visceral pathways. *Development* **130**:6635–6642. doi: [10.1242/dev.00866](https://doi.org/10.1242/dev.00866), PMID: [14627719](https://pubmed.ncbi.nlm.nih.gov/14627719/)
- Dunel-Erb S**, Bailly Y, Laurent P. 1982. Neuroepithelial cells in fish gill primary lamellae. *Journal of Applied Physiology* **53**:1342–1353. PMID: [7153134](https://pubmed.ncbi.nlm.nih.gov/7153134/)
- Engert S**, Liao WP, Bartscher I, Lickert H. 2009. Sox17-2A-iCre: a knock-in mouse line expressing cre recombinase in endoderm and vascular endothelial cells. *Genesis* **47**:603–610. doi: [10.1002/dvg.20540](https://doi.org/10.1002/dvg.20540), PMID: [19548312](https://pubmed.ncbi.nlm.nih.gov/19548312/)
- Freem LJ**, Escot S, Tannahill D, Druckenbrod NR, Thapar N, Burns AJ. 2010. The intrinsic innervation of the lung is derived from neural crest cells as shown by optical projection tomography in *Wnt1-Cre;YFP* reporter mice. *Journal of Anatomy* **217**:651–664. doi: [10.1111/j.1469-7580.2010.01295.x](https://doi.org/10.1111/j.1469-7580.2010.01295.x), PMID: [20840354](https://pubmed.ncbi.nlm.nih.gov/20840354/)
- Gage PJ**, Rhoades W, Prucka SK, Hjalt T. 2005. Fate maps of neural crest and mesoderm in the mammalian eye. *Investigative Ophthalmology & Visual Science* **46**:4200–4208. doi: [10.1167/iovs.05-0691](https://doi.org/10.1167/iovs.05-0691), PMID: [16249499](https://pubmed.ncbi.nlm.nih.gov/16249499/)
- García-Fernández M**, Mejías R, López-Barneo J. 2007. Developmental changes of chromaffin cell secretory response to hypoxia studied in thin adrenal slices. *Pflügers Archiv - European Journal of Physiology* **454**:93–100. doi: [10.1007/s00424-006-0186-y](https://doi.org/10.1007/s00424-006-0186-y), PMID: [17165070](https://pubmed.ncbi.nlm.nih.gov/17165070/)
- Gaskell JF**. 1912. The distribution and physiological action of the suprarenal medullary tissue in *Petromyzon fluviatilis*. *The Journal of Physiology* **44**:59–67. doi: [10.1113/jphysiol.1912.sp001502](https://doi.org/10.1113/jphysiol.1912.sp001502), PMID: [16993137](https://pubmed.ncbi.nlm.nih.gov/16993137/)
- Giacomini E**. 1902. Contributo alla conoscenza delle capsule surrenali dei Ciclostomi. Sulle capsule surrenali dei Petromizonti. *Monitore Zoologico Italiano* **13**:143–162.
- Gillis JA**, Tidswell ORA. 2017. The origin of vertebrate gills. *Current Biology* **27**:729–732. doi: [10.1016/j.cub.2017.01.022](https://doi.org/10.1016/j.cub.2017.01.022), PMID: [28190727](https://pubmed.ncbi.nlm.nih.gov/28190727/)
- Guha A**, Vasconcelos M, Cai Y, Yoneda M, Hinds A, Qian J, Li G, Dickel L, Johnson JE, Kimura S, Guo J, McMahon J, McMahon AP, Cardoso WV. 2012. Neuroepithelial body microenvironment is a niche for a distinct subset of Clara-like precursors in the developing airways. *PNAS* **109**:12592–12597. doi: [10.1073/pnas.1204710109](https://doi.org/10.1073/pnas.1204710109), PMID: [22797898](https://pubmed.ncbi.nlm.nih.gov/22797898/)
- Häming D**, Simoes-Costa M, Uy B, Valencia J, Sauka-Spengler T, Bronner-Fraser M. 2011. Expression of sympathetic nervous system genes in lamprey suggests their recruitment for specification of a new vertebrate feature. *PLoS One* **6**:e26543. doi: [10.1371/journal.pone.0026543](https://doi.org/10.1371/journal.pone.0026543), PMID: [22046306](https://pubmed.ncbi.nlm.nih.gov/22046306/)
- Hempleman SC**, Warburton SJ. 2013. Comparative embryology of the carotid body. *Respiratory Physiology & Neurobiology* **185**:3–8. doi: [10.1016/j.resp.2012.08.004](https://doi.org/10.1016/j.resp.2012.08.004), PMID: [22902512](https://pubmed.ncbi.nlm.nih.gov/22902512/)

- Hoyt RF, Jr.,** McNelly NA, Sorokin SP. 1990. Dynamics of neuroepithelial body (NEB) formation in developing hamster lung: light microscopic autoradiography after 3H-thymidine labeling in vivo. *The Anatomical Record* **227**:340–350. doi: [10.1002/ar.1092270309](https://doi.org/10.1002/ar.1092270309), PMID: [2372138](https://pubmed.ncbi.nlm.nih.gov/2372138/)
- Hsia CCW,** Schmitz A, Lambert M, Perry SF, Maina JN. 2013. Evolution of air breathing: oxygen homeostasis and the transitions from water to land and sky. *Comprehensive Physiology* **3**:849–915. doi: [10.1002/cphy.c120003](https://doi.org/10.1002/cphy.c120003), PMID: [23720333](https://pubmed.ncbi.nlm.nih.gov/23720333/)
- Hu W,** Haamedi N, Lee J, Kinoshita T, Ohnuma S. 2013. The structure and development of *Xenopus laevis* cornea. *Experimental Eye Research* **116**:109–128. doi: [10.1016/j.exer.2013.07.021](https://doi.org/10.1016/j.exer.2013.07.021), PMID: [23896054](https://pubmed.ncbi.nlm.nih.gov/23896054/)
- Ishii K,** Honda K, Ishii K. 1966. The function of the carotid labyrinth in the toad. *The Tohoku Journal of Experimental Medicine* **88**:103–116. doi: [10.1620/tjem.88.103](https://doi.org/10.1620/tjem.88.103), PMID: [5955235](https://pubmed.ncbi.nlm.nih.gov/5955235/)
- Ito T,** Udaka N, Yazawa T, Okudela K, Hayashi H, Sudo T, Guillemot F, Kageyama R, Kitamura H. 2000. Basic helix-loop-helix transcription factors regulate the neuroendocrine differentiation of fetal mouse pulmonary epithelium. *Development* **127**:3913–3921. PMID: [10952889](https://pubmed.ncbi.nlm.nih.gov/10952889/)
- Johnston MC,** Noden DM, Hazelton RD, Coulombre JL, Coulombre AJ. 1979. Origins of avian ocular and periocular tissues. *Experimental Eye Research* **29**:27–43. doi: [10.1016/0014-4835\(79\)90164-7](https://doi.org/10.1016/0014-4835(79)90164-7)
- Jonz MG,** Nurse CA. 2003. Neuroepithelial cells and associated innervation of the zebrafish gill: a confocal immunofluorescence study. *The Journal of Comparative Neurology* **461**:1–17. doi: [10.1002/cne.10680](https://doi.org/10.1002/cne.10680), PMID: [12722101](https://pubmed.ncbi.nlm.nih.gov/12722101/)
- Jonz MG,** Nurse CA. 2005. Development of oxygen sensing in the gills of zebrafish. *Journal of Experimental Biology* **208**:1537–1549. doi: [10.1242/jeb.01564](https://doi.org/10.1242/jeb.01564), PMID: [15802677](https://pubmed.ncbi.nlm.nih.gov/15802677/)
- Jonz MG,** Nurse CA. 2006. Ontogenesis of oxygen chemoreception in aquatic vertebrates. *Respiratory Physiology & Neurobiology* **154**:139–152. doi: [10.1016/j.resp.2006.01.004](https://doi.org/10.1016/j.resp.2006.01.004), PMID: [16488670](https://pubmed.ncbi.nlm.nih.gov/16488670/)
- Jonz MG,** Fearon IM, Nurse CA. 2004. Neuroepithelial oxygen chemoreceptors of the zebrafish gill. *The Journal of Physiology* **560**:737–752. doi: [10.1113/jphysiol.2004.069294](https://doi.org/10.1113/jphysiol.2004.069294), PMID: [15331683](https://pubmed.ncbi.nlm.nih.gov/15331683/)
- Jonz MG,** Buck LT, Perry SF, Schwerte T, Zaccane G. 2016. Sensing and surviving hypoxia in vertebrates. *Annals of the New York Academy of Sciences* **1365**:43–58. doi: [10.1111/nyas.12780](https://doi.org/10.1111/nyas.12780), PMID: [25959851](https://pubmed.ncbi.nlm.nih.gov/25959851/)
- Kague E,** Gallagher M, Burke S, Parsons M, Franz-Odenaal T, Fisher S. 2012. Skeletogenic fate of zebrafish cranial and trunk neural crest. *PLoS One* **7**:e47394. doi: [10.1371/journal.pone.0047394](https://doi.org/10.1371/journal.pone.0047394), PMID: [23155370](https://pubmed.ncbi.nlm.nih.gov/23155370/)
- Kameda Y.** 2002. Carotid body and glomus cells distributed in the wall of the common carotid artery in the bird. *Microscopy Research and Technique* **59**:196–206. doi: [10.1002/jemt.10194](https://doi.org/10.1002/jemt.10194), PMID: [12384964](https://pubmed.ncbi.nlm.nih.gov/12384964/)
- Kameda Y.** 2005. *Mash1* is required for glomus cell formation in the mouse carotid body. *Developmental Biology* **283**:128–139. doi: [10.1016/j.ydbio.2005.04.004](https://doi.org/10.1016/j.ydbio.2005.04.004), PMID: [15878769](https://pubmed.ncbi.nlm.nih.gov/15878769/)
- Kaufman CK,** Mosimann C, Fan ZP, Yang S, Thomas AJ, Ablain J, Tan JL, Fogley RD, van Rooijen E, Hagedorn EJ, Ciarlo C, White RM, Matos DA, Puller AC, Santoriello C, Liao EC, Young RA, Zon LI. 2016. A zebrafish melanoma model reveals emergence of neural crest identity during melanoma initiation. *Science* **351**:aad2197. doi: [10.1126/science.aad2197](https://doi.org/10.1126/science.aad2197), PMID: [26823433](https://pubmed.ncbi.nlm.nih.gov/26823433/)
- Kuo CS,** Krasnow MA. 2015. Formation of a neurosensory organ by epithelial cell slithering. *Cell* **163**:394–405. doi: [10.1016/j.cell.2015.09.021](https://doi.org/10.1016/j.cell.2015.09.021), PMID: [26435104](https://pubmed.ncbi.nlm.nih.gov/26435104/)
- Kusakabe T.** 2002. Carotid labyrinth of amphibians. *Microscopy Research and Technique* **59**:207–226. doi: [10.1002/jemt.10195](https://doi.org/10.1002/jemt.10195), PMID: [12384965](https://pubmed.ncbi.nlm.nih.gov/12384965/)
- Kusakabe T.** 2009. The amphibian carotid labyrinth. In: Zaccane G, Cutz E, Adriaensen D, Nurse CA, Mauceri A (Eds). *Airway Chemoreceptors in the Vertebrates: Structure, Evolution and Function*. Enfield, New Hampshire, USA: Science Publishers. p. 125–139.
- Kuscha V,** Barreiro-Iglesias A, Becker CG, Becker T. 2012. Plasticity of tyrosine hydroxylase and serotonergic systems in the regenerating spinal cord of adult zebrafish. *The Journal of Comparative Neurology* **520**:933–951. doi: [10.1002/cne.22739](https://doi.org/10.1002/cne.22739), PMID: [21830219](https://pubmed.ncbi.nlm.nih.gov/21830219/)
- Le Douarin N,** Le Lièvre C, Fontaine J. 1972. Recherches expérimentales sur l'origine embryologique du corps carotidien chez les Oiseaux. *Comptes Rendus De l'Académie Des Sciences* **275**:583–586.
- Le Douarin NM.** 2004. The avian embryo as a model to study the development of the neural crest: a long and still ongoing story. *Mechanisms of Development* **121**:1089–1102. doi: [10.1016/j.mod.2004.06.003](https://doi.org/10.1016/j.mod.2004.06.003), PMID: [15296974](https://pubmed.ncbi.nlm.nih.gov/15296974/)
- Levitsky KL,** López-Barneo J. 2009. Developmental change of T-type Ca^{2+} channel expression and its role in rat chromaffin cell responsiveness to acute hypoxia. *The Journal of Physiology* **587**:1917–1929. doi: [10.1113/jphysiol.2009.168989](https://doi.org/10.1113/jphysiol.2009.168989), PMID: [19273573](https://pubmed.ncbi.nlm.nih.gov/19273573/)
- Lo L-C,** Johnson JE, Wuenschell CW, Saito T, Anderson DJ. 1991. Mammalian *achaete-scute* homolog 1 is transiently expressed by spatially restricted subsets of early neuroepithelial and neural crest cells. *Genes & Development* **5**:1524–1537. doi: [10.1101/gad.5.9.1524](https://doi.org/10.1101/gad.5.9.1524), PMID: [1909283](https://pubmed.ncbi.nlm.nih.gov/1909283/)
- López-Barneo J,** López-López JR, Ureña J, González C. 1988. Chemotransduction in the carotid body: K^+ current modulated by PO_2 in type I chemoreceptor cells. *Science* **241**:580–582. doi: [10.1126/science.2456613](https://doi.org/10.1126/science.2456613), PMID: [2456613](https://pubmed.ncbi.nlm.nih.gov/2456613/)
- López-Barneo J,** Ortega-Sáenz P, González-Rodríguez P, Fernández-Agüera MC, Macías D, Pardo R, Gao L. 2016. Oxygen-sensing by arterial chemoreceptors: mechanisms and medical translation. *Molecular Aspects of Medicine* **47-48**:90–108. doi: [10.1016/j.mam.2015.12.002](https://doi.org/10.1016/j.mam.2015.12.002), PMID: [26709054](https://pubmed.ncbi.nlm.nih.gov/26709054/)
- Madisen L,** Zwingman TA, Sunkin SM, Oh SW, Zariwala HA, Gu H, Ng LL, Palmiter RD, Hawrylycz MJ, Jones AR, Lein ES, Zeng H. 2010. A robust and high-throughput Cre reporting and characterization system for the whole mouse brain. *Nature Neuroscience* **13**:133–140. doi: [10.1038/nn.2467](https://doi.org/10.1038/nn.2467), PMID: [20023653](https://pubmed.ncbi.nlm.nih.gov/20023653/)

- Marsh-Armstrong N**, Huang H, Berry DL, Brown DD. 1999. Germ-line transmission of transgenes in *Xenopus laevis*. *PNAS* **96**:14389–14393. doi: [10.1073/pnas.96.25.14389](https://doi.org/10.1073/pnas.96.25.14389), PMID: [10588715](https://pubmed.ncbi.nlm.nih.gov/10588715/)
- McCauley DW**, Bronner-Fraser M. 2003. Neural crest contributions to the lamprey head. *Development* **130**: 2317–2327. doi: [10.1242/dev.00451](https://doi.org/10.1242/dev.00451), PMID: [12702647](https://pubmed.ncbi.nlm.nih.gov/12702647/)
- McGrew MJ**, Sherman A, Lillico SG, Ellard FM, Radcliffe PA, Gilhooley HJ, Mitrophanous KA, Cambray N, Wilson V, Sang H. 2008. Localised axial progenitor cell populations in the avian tail bud are not committed to a posterior hox identity. *Development* **135**:2289–2299. doi: [10.1242/dev.022020](https://doi.org/10.1242/dev.022020), PMID: [18508860](https://pubmed.ncbi.nlm.nih.gov/18508860/)
- Metcalfe WK**, Myers PZ, Trevarrow B, Bass MB, Kimmel CB. 1990. Primary neurons that express the L2/HNK-1 carbohydrate during early development in the zebrafish. *Development* **110**:491–504. PMID: [1723944](https://pubmed.ncbi.nlm.nih.gov/1723944/)
- Miller SR**, Perera SN, Baker CVH. 2017. Constitutively active Notch1 converts cranial neural crest-derived frontonasal mesenchyme to perivascular cells *in vivo*. *Biology Open* **6**:317–325. doi: [10.1242/bio.023887](https://doi.org/10.1242/bio.023887), PMID: [28183698](https://pubmed.ncbi.nlm.nih.gov/28183698/)
- Milsom WK**. 2012. New insights into gill chemoreception: receptor distribution and roles in water and air breathing fish. *Respiratory Physiology & Neurobiology* **184**:326–339. doi: [10.1016/j.resp.2012.07.013](https://doi.org/10.1016/j.resp.2012.07.013), PMID: [22841952](https://pubmed.ncbi.nlm.nih.gov/22841952/)
- Milsom WK**, Bursleson ML. 2007. Peripheral arterial chemoreceptors and the evolution of the carotid body. *Respiratory Physiology & Neurobiology* **157**:4–11. doi: [10.1016/j.resp.2007.02.007](https://doi.org/10.1016/j.resp.2007.02.007), PMID: [17353155](https://pubmed.ncbi.nlm.nih.gov/17353155/)
- Mizoguchi T**, Verkade H, Heath JK, Kuroiwa A, Kikuchi Y. 2008. Sdf1/Cxcr4 signaling controls the dorsal migration of endodermal cells during zebrafish gastrulation. *Development* **135**:2521–2529. doi: [10.1242/dev.020107](https://doi.org/10.1242/dev.020107), PMID: [18579679](https://pubmed.ncbi.nlm.nih.gov/18579679/)
- Mongera A**, Singh AP, Levesque MP, Chen Y-Y, Konstantinidis P, Nüsslein-Volhard C. 2013. Genetic lineage labeling in zebrafish uncovers novel neural crest contributions to the head, including gill pillar cells. *Development* **140**:916–925. doi: [10.1242/dev.091066](https://doi.org/10.1242/dev.091066), PMID: [23362350](https://pubmed.ncbi.nlm.nih.gov/23362350/)
- Montero-Balaguer M**, Lang MR, Sachdev SW, Knappmeyer C, Stewart RA, De La Guardia A, Hatzopoulos AK, Knapik EW. 2006. The mother superior mutation ablates *foxd3* activity in neural crest progenitor cells and depletes neural crest derivatives in zebrafish. *Developmental Dynamics* **235**:3199–3212. doi: [10.1002/dvdy.20959](https://doi.org/10.1002/dvdy.20959), PMID: [17013879](https://pubmed.ncbi.nlm.nih.gov/17013879/)
- Mosimann C**, Kaufman CK, Li P, Pugach EK, Tamplin OJ, Zon LI. 2011. Ubiquitous transgene expression and Cre-based recombination driven by the ubiquitin promoter in zebrafish. *Development* **138**:169–177. doi: [10.1242/dev.059345](https://doi.org/10.1242/dev.059345), PMID: [21138979](https://pubmed.ncbi.nlm.nih.gov/21138979/)
- Nieuwkoop PD**, Faber J. 1967. *Normal Table of Xenopus laevis (Daudin)*. Amsterdam: North-Holland.
- Nikitina N**, Bronner-Fraser M, Sauka-Spengler T. 2009. The sea lamprey *Petromyzon marinus*: a model for evolutionary and developmental biology. In: *Emerging Model Organisms: A Laboratory Manual*. New York: CSHL Press. p. 405–429.
- Noden DM**. 1978. The control of avian cephalic neural crest cytodifferentiation. I. skeletal and connective tissues. *Developmental Biology* **67**:296–312. doi: [10.1016/0012-1606\(78\)90201-4](https://doi.org/10.1016/0012-1606(78)90201-4), PMID: [738529](https://pubmed.ncbi.nlm.nih.gov/738529/)
- Nurse CA**, Piskuric NA. 2013. Signal processing at mammalian carotid body chemoreceptors. *Seminars in Cell & Developmental Biology* **24**:22–30. doi: [10.1016/j.semcdb.2012.09.006](https://doi.org/10.1016/j.semcdb.2012.09.006), PMID: [23022231](https://pubmed.ncbi.nlm.nih.gov/23022231/)
- Nurse CA**. 2014. Synaptic and paracrine mechanisms at carotid body arterial chemoreceptors. *The Journal of Physiology* **592**:3419–3426. doi: [10.1113/jphysiol.2013.269829](https://doi.org/10.1113/jphysiol.2013.269829), PMID: [24665097](https://pubmed.ncbi.nlm.nih.gov/24665097/)
- Pardal R**, Ortega-Sáenz P, Durán R, López-Barneo J. 2007. Glia-like stem cells sustain physiologic neurogenesis in the adult mammalian carotid body. *Cell* **131**:364–377. doi: [10.1016/j.cell.2007.07.043](https://doi.org/10.1016/j.cell.2007.07.043), PMID: [17956736](https://pubmed.ncbi.nlm.nih.gov/17956736/)
- Park K-S**, Liang M-C, Raiser DM, Zamponi R, Roach RR, Curtis SJ, Walton Z, Schaffer BE, Roake CM, Zmoos A-F, Kriegel C, Wong K-K, Sage J, Kim CF. 2011. Characterization of the cell of origin for small cell lung cancer. *Cell Cycle* **10**:2806–2815. doi: [10.4161/cc.10.16.17012](https://doi.org/10.4161/cc.10.16.17012), PMID: [21822053](https://pubmed.ncbi.nlm.nih.gov/21822053/)
- Pearse AG**, Polak JM, Rost FW, Fontaine J, Le Lièvre C, Le Douarin N. 1973. Demonstration of the neural crest origin of type I (APUD) cells in the avian carotid body, using a cytochemical marker system. *Histochemie* **34**: 191–203. doi: [10.1007/BF00303435](https://doi.org/10.1007/BF00303435), PMID: [4693636](https://pubmed.ncbi.nlm.nih.gov/4693636/)
- Porteus CS**, Brink DL, Coolidge EH, Fong AY, Milsom WK. 2013. Distribution of acetylcholine and catecholamines in fish gills and their potential roles in the hypoxic ventilatory response. *Acta Histochemica* **115**:158–169. doi: [10.1016/j.acthis.2012.06.004](https://doi.org/10.1016/j.acthis.2012.06.004), PMID: [22765871](https://pubmed.ncbi.nlm.nih.gov/22765871/)
- Porteus CS**, Wright PA, Milsom WK. 2014. Characterisation of putative oxygen chemoreceptors in bowfin (*Amia calva*). *Journal of Experimental Biology* **217**:1269–1277. doi: [10.1242/jeb.098467](https://doi.org/10.1242/jeb.098467), PMID: [24744423](https://pubmed.ncbi.nlm.nih.gov/24744423/)
- Qin Z**, Lewis JE, Perry SF. 2010. Zebrafish (*Danio rerio*) gill neuroepithelial cells are sensitive chemoreceptors for environmental CO₂. *The Journal of Physiology* **588**:861–872. doi: [10.1113/jphysiol.2009.184739](https://doi.org/10.1113/jphysiol.2009.184739), PMID: [20051495](https://pubmed.ncbi.nlm.nih.gov/20051495/)
- Rawlins EL**, Clark CP, Xue Y, Hogan BL. 2009. The Id2⁺ distal tip lung epithelium contains individual multipotent embryonic progenitor cells. *Development* **136**:3741–3745. doi: [10.1242/dev.037317](https://doi.org/10.1242/dev.037317), PMID: [19855016](https://pubmed.ncbi.nlm.nih.gov/19855016/)
- Regan KS**, Jonz MG, Wright PA. 2011. Neuroepithelial cells and the hypoxia emersion response in the amphibious fish *Kryptolebias marmoratus*. *Journal of Experimental Biology* **214**:2560–2568. doi: [10.1242/jeb.056333](https://doi.org/10.1242/jeb.056333), PMID: [21753050](https://pubmed.ncbi.nlm.nih.gov/21753050/)
- Reyes C**, Fong AY, Brink DL, Milsom WK. 2014. Distribution and innervation of putative arterial chemoreceptors in the bullfrog (*Rana catesbeiana*). *Journal of Comparative Neurology* **522**:3754–3774. doi: [10.1002/cne.23640](https://doi.org/10.1002/cne.23640), PMID: [24954002](https://pubmed.ncbi.nlm.nih.gov/24954002/)
- Reynolds SD**, Giangreco A, Power JHT, Stripp BR. 2000. Neuroepithelial bodies of pulmonary airways serve as a reservoir of progenitor cells capable of epithelial regeneration. *The American Journal of Pathology* **156**:269–278. doi: [10.1016/S0002-9440\(10\)64727-X](https://doi.org/10.1016/S0002-9440(10)64727-X), PMID: [10623675](https://pubmed.ncbi.nlm.nih.gov/10623675/)

- Ruppert EE**, Fox RS, Barnes RB. 2003. *Invertebrate Zoology: A Functional Evolutionary Approach*. Belmont, California, United States: Brooks/Cole Thompson Learning.
- Sadaghiani B**, Thiébaud CH. 1987. Neural crest development in the *Xenopus laevis* embryo, studied by interspecific transplantation and scanning electron microscopy. *Developmental Biology* **124**:91–110. doi: [10.1016/0012-1606\(87\)90463-5](https://doi.org/10.1016/0012-1606(87)90463-5), PMID: [3666314](https://pubmed.ncbi.nlm.nih.gov/3666314/)
- Saltys HA**, Jonz MG, Nurse CA. 2006. Comparative study of gill neuroepithelial cells and their innervation in teleosts and *Xenopus* tadpoles. *Cell and Tissue Research* **323**:1–10. doi: [10.1007/s00441-005-0048-5](https://doi.org/10.1007/s00441-005-0048-5), PMID: [16163489](https://pubmed.ncbi.nlm.nih.gov/16163489/)
- Schlosser G**. 2003. Hypobranchial placodes in *Xenopus laevis* give rise to hypobranchial ganglia, a novel type of cranial ganglia. *Cell and Tissue Research* **312**:21–29. doi: [10.1007/s00441-003-0710-8](https://doi.org/10.1007/s00441-003-0710-8), PMID: [12712315](https://pubmed.ncbi.nlm.nih.gov/12712315/)
- Seidler FJ**, Slotkin TA. 1985. Adrenomedullary function in the neonatal rat: responses to acute hypoxia. *The Journal of Physiology* **358**:1–16. doi: [10.1113/jphysiol.1985.sp015536](https://doi.org/10.1113/jphysiol.1985.sp015536), PMID: [2858585](https://pubmed.ncbi.nlm.nih.gov/2858585/)
- Song H**, Yao E, Lin C, Gacayan R, Chen M-H, Chuang P-T. 2012. Functional characterization of pulmonary neuroendocrine cells in lung development, injury, and tumorigenesis. *PNAS* **109**:17531–17536. doi: [10.1073/pnas.1207238109](https://doi.org/10.1073/pnas.1207238109), PMID: [23047698](https://pubmed.ncbi.nlm.nih.gov/23047698/)
- Soriano P**. 1999. Generalized lacZ expression with the ROSA26 Cre reporter strain. *Nature Genetics* **21**:70–71. doi: [10.1038/5007](https://doi.org/10.1038/5007), PMID: [9916792](https://pubmed.ncbi.nlm.nih.gov/9916792/)
- Soukup V**, Epperlein H-H, Horáček I, Cerný R. 2008. Dual epithelial origin of vertebrate oral teeth. *Nature* **455**:795–798. doi: [10.1038/nature07304](https://doi.org/10.1038/nature07304), PMID: [18794902](https://pubmed.ncbi.nlm.nih.gov/18794902/)
- Soules KA**, Link BA. 2005. Morphogenesis of the anterior segment in the zebrafish eye. *BMC Developmental Biology* **5**:12. doi: [10.1186/1471-213X-5-12](https://doi.org/10.1186/1471-213X-5-12), PMID: [15985175](https://pubmed.ncbi.nlm.nih.gov/15985175/)
- Srinivas S**, Watanabe T, Lin C-S, William CM, Tanabe Y, Jessell TM, Costantini F. 2001. Cre reporter strains produced by targeted insertion of EYFP and ECFP into the ROSA26 locus. *BMC Developmental Biology* **1**:4. doi: [10.1186/1471-213X-1-4](https://doi.org/10.1186/1471-213X-1-4), PMID: [11299042](https://pubmed.ncbi.nlm.nih.gov/11299042/)
- Stanke M**, Junghans D, Geissen M, Goridis C, Ernsberger U, Rohrer H. 1999. The Phox2 homeodomain proteins are sufficient to promote the development of sympathetic neurons. *Development* **126**:4087–4094. PMID: [10457017](https://pubmed.ncbi.nlm.nih.gov/10457017/)
- Stone LM**, Finger TE, Tam PPL, Tan S-S. 1995. Taste receptor cells arise from local epithelium, not neurogenic ectoderm. *PNAS* **92**:1916–1920. doi: [10.1073/pnas.92.6.1916](https://doi.org/10.1073/pnas.92.6.1916), PMID: [7892199](https://pubmed.ncbi.nlm.nih.gov/7892199/)
- Sundvik M**, Chen Y-C, Panula P. 2013. Presenilin1 regulates histamine neuron development and behavior in zebrafish, *Danio rerio*. *Journal of Neuroscience* **33**:1589–1597. doi: [10.1523/JNEUROSCI.1802-12.2013](https://doi.org/10.1523/JNEUROSCI.1802-12.2013), PMID: [23345232](https://pubmed.ncbi.nlm.nih.gov/23345232/)
- Sutherland KD**, Proost N, Brouns I, Adriaensen D, Song J-Y, Berns A. 2011. Cell of origin of small cell lung cancer: inactivation of *Trp53* and *Rb1* in distinct cell types of adult mouse lung. *Cancer Cell* **19**:754–764. doi: [10.1016/j.ccr.2011.04.019](https://doi.org/10.1016/j.ccr.2011.04.019), PMID: [21665149](https://pubmed.ncbi.nlm.nih.gov/21665149/)
- Suzuki DG**, Murakami Y, Escriva H, Wada H. 2015. A comparative examination of neural circuit and brain patterning between the lamprey and amphioxus reveals the evolutionary origin of the vertebrate visual center. *Journal of Comparative Neurology* **523**:251–261. doi: [10.1002/cne.23679](https://doi.org/10.1002/cne.23679), PMID: [25233869](https://pubmed.ncbi.nlm.nih.gov/25233869/)
- Tahara Y**. 1988. Normal stages of development in the lamprey *Lampetra reissneri* (Dybowski). *Zoological Science* **5**:109–118.
- Teppema LJ**, Dahan A. 2010. The ventilatory response to hypoxia in mammals: mechanisms, measurement, and analysis. *Physiological Reviews* **90**:675–754. doi: [10.1152/physrev.00012.2009](https://doi.org/10.1152/physrev.00012.2009), PMID: [20393196](https://pubmed.ncbi.nlm.nih.gov/20393196/)
- Thompson RJ**, Jackson A, Nurse CA. 1997. Developmental loss of hypoxic chemosensitivity in rat adrenomedullary chromaffin cells. *The Journal of Physiology* **498**:503–510. doi: [10.1113/jphysiol.1997.sp021876](https://doi.org/10.1113/jphysiol.1997.sp021876), PMID: [9032697](https://pubmed.ncbi.nlm.nih.gov/9032697/)
- Voshol H**, van Zuylen CWEM, Orberger G, Vliegenthart JFG, Schachner M. 1996. Structure of the HNK-1 carbohydrate epitope on bovine peripheral myelin glycoprotein P0. *The Journal of Biological Chemistry* **271**:22957–22960. doi: [10.1074/jbc.271.38.22957](https://doi.org/10.1074/jbc.271.38.22957), PMID: [8798480](https://pubmed.ncbi.nlm.nih.gov/8798480/)
- Wang W-D**, Melville DB, Montero-Balaguer M, Hatzopoulos AK, Knapik EW. 2011. Tfp2a and Foxd3 regulate early steps in the development of the neural crest progenitor population. *Developmental Biology* **360**:173–185. doi: [10.1016/j.ydbio.2011.09.019](https://doi.org/10.1016/j.ydbio.2011.09.019), PMID: [21963426](https://pubmed.ncbi.nlm.nih.gov/21963426/)
- Warga RM**, Nüsslein-Volhard C. 1999. Origin and development of the zebrafish endoderm. *Development* **126**:827–838. PMID: [9895329](https://pubmed.ncbi.nlm.nih.gov/9895329/)
- Yamamoto K**, Ruuskanen JO, Wullmann MF, Vernier P. 2011. Differential expression of dopaminergic cell markers in the adult zebrafish forebrain. *The Journal of Comparative Neurology* **519**:576–598. doi: [10.1002/cne.22535](https://doi.org/10.1002/cne.22535), PMID: [21192085](https://pubmed.ncbi.nlm.nih.gov/21192085/)
- Youngson C**, Nurse C, Yeager H, Cutz E. 1993. Oxygen sensing in airway chemoreceptors. *Nature* **365**:153–155. doi: [10.1038/365153a0](https://doi.org/10.1038/365153a0), PMID: [8371757](https://pubmed.ncbi.nlm.nih.gov/8371757/)
- Yuan G**, Peng Y-J, Reddy VD, Makarenko VV, Nanduri J, Khan SA, Garcia JA, Kumar GK, Semenza GL, Prabhakar NR. 2013. Mutual antagonism between hypoxia-inducible factors 1 α and 2 α regulates oxygen sensing and cardio-respiratory homeostasis. *PNAS* **110**:E1788–E1796. doi: [10.1073/pnas.1305961110](https://doi.org/10.1073/pnas.1305961110), PMID: [23610397](https://pubmed.ncbi.nlm.nih.gov/23610397/)
- Zhao XC**, Yee RW, Norcom E, Burgess H, Avanesov AS, Barrish JP, Malicki J. 2006. The zebrafish cornea: structure and development. *Investigative Ophthalmology & Visual Science* **47**:4341–4348. doi: [10.1167/iovs.05-1611](https://doi.org/10.1167/iovs.05-1611), PMID: [17003424](https://pubmed.ncbi.nlm.nih.gov/17003424/)